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TITLE OF THE INVENTION

Method for Normalizing the Relative Intensities of Detection Signals in Hybridization Arrays

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FIELD OF THE INVENTION

The present invention relates to the field of hybridization arrays. More specifically, the present invention concerns a method for normalizing signals to be compared in hybridization arrays. This novel method relies on the use of ribosomal RNA (rRNA) as an internal standard and allows approximation of the relative abundance of multiple mRNAs as well as direct comparisons between any two specific RNA samples.

BACKGROUND OF THE INVENTION

In DNA microarray experiments, one of the more popular ways to control for spotted DNA quantity and surface chemistry anomalies involves the use of two-color fluorescence (see refs. 4, 5). For example, a Cy3 (green)-labelled probe prepared from healthy tissue could be used as a control to examine expression profiles of a Cy5 (red)-labelled probe prepared from a tumor tissue. The normalized expression values for every gene would then be calculated as the ratio of experimental expression to control expression. This method can obviously eliminate much (but not all) experimental variation by allowing two samples to be compared on the same chip because there is enough DNA on each spot that both test and reference cDNAs can hybridize to it at once without interference. More sophisticated three-color experiments are also possible in which one channel serves as a control for the amount of spotted DNA, and channels two and three allow two samples to be compared to this control and to each other (see ref. 5).

In addition to the local normalization method described above, more general methods are also available in the form of control spots on the slide. With a set of control spots, it is possible to control variations in overall slide quality or scanning differences.

Applicable normalization strategies are based on some underlying assumptions regarding the data and the strategies used for each experiment. These

strategies must therefore be adjusted to reflect both the system under study and the experimental design. A primary assumption is that for either an entire collection of arrayed genes or some subset of the genes (such as housekeeping genes), or for some added set of controls, the ratio of measured expression averaged over the set should be close to unity.

The need for good methods of normalisation for microarray data can not be overstated (see refs. 6, 7). Depending on the experimental design, there are three useful approaches for calculating normalization factors. The first simply relies on the total fluorescent intensity measured. The assumption underlying this approach is that the total mass of RNA labelled with either Cy3 or Cy5 is equal. While the intensity for any one spot may be higher in one channel than the other, when averaged over thousands of spots in a given array, these fluctuations average out. Consequently, the total integrated intensity across all the spots in the array should be equal for both channels. Alternatively, one could add a number of controls in increasing but equimolar concentrations to both labeling reactions, and the sum of the intensities for these spots should be equal.

A second approach uses linear regression analysis. For closely related samples, one would expect many of the genes to be expressed at nearly constant levels. Consequently, a scatter plot of the measured Cy5 versus Cy3 intensities should have a slope of one. Measured intensities for added equimolar controls should behave similarly. Under this assumption, one can use regression analysis techniques to calculate the slope which is used to rescale the data and adjust the slope to one.

A third approach has been described by Chen et al (1997) (ref. 1). In it, it is assumed that a subset of housekeeping genes exists and that for these genes the distribution of transcription levels should have some mean value and standard deviation that are independent of any particular sample. In this case, the ratio of measured Cy5 to Cy3 ratios for these genes can be modeled and the mean of the ratio adjusted to 1. Chen and his collaborators describe an iterative procedure to achieve this normalization. Quackenbush and collaborators (ref. 2) have implemented their own algorithm and a variation thereof that uses the entire data set in a data visualization and analysis tool called TIGR ArrayViewer. Other statistical methods of determining data accuracy have been described (ref. 3, 11).

The above procedures describe array-based measures that can be used to normalize data. However, even with multiple colour fluorescence and control spots, undesired experimental variation can contaminate expression data. It is also possible that some or all of the physical normalization techniques are
5 missing from the experiment, in which case it is even more important to find additional means of normalization.

The use of internal standards overcomes these problems. Using an exogenously added standard has the advantage of giving the user absolute
10 control over the amount of template added, with no variation between samples. Using an exogenous standard does not, however, control differences in the quality of the starting RNA in a reverse transcription reaction. If there are differences in the levels of integrity of the RNA between otherwise identical samples, the yield of specific reverse transcriptase products will reflect this
15 variation, although the external standards will still appear identical. For this reason, as well as for simplicity and reproducibility, an endogenous RNA standard should be favoured in microarray experiments.

Theoretically, an ideal endogenous standard for a DNA microarray would be a
20 transcript whose expression does not vary during the cell cycle, between cell types, or in response to the experimental treatments that one wishes to examine. Additionally, for an endogenous standard to be valid in a microarray it is crucial that it be of a similar relative abundance as the test and reference (or target) transcripts in the microarray. Unfortunately, such a molecule does
25 not exist and there are serious limitations to the standards currently in use. For example, although beta-actin is a frequently used standard (refs 9, 10), its level of expression varies significantly from tissue to tissue.

For DNA microarray experiments, mRNA is copied into cDNA with the use of
30 reverse transcriptase so that the relative abundance of individual mRNAs is reflected in the cDNA product. Input RNA in reverse transcription reactions is usually quantified by spectrophotometry. The RNA that is used in a typical pre-reverse transcription reaction is total RNA, 80% of which is ribosomal RNA. The mRNA component of total cellular RNA can vary from 2% to 5% depending on
35 the tissue, the remainder of the RNA consisting of tRNA or small nuclear RNAs. Therefore, even if a transcript is invariant (as expressed as a percentage of mRNA), its relative abundance would still vary when considered as a percent of the total input RNA from different source tissues.

Since the majority of the RNA is rRNA, the level of rRNA remains essentially constant from sample to sample. Because 18S and 28S rRNA make up the majority of optically absorbent material at OD_{260nm}, they should make ideal invariant controls. In fact, 18S and 28S transcripts are frequently used as
5 internal controls in northern hybridization, RNase protection and quantitative RT-PCR assays (see ref. 8). However, the overwhelming abundance of rRNA is a major limitation to its utility as a control in DNA microarray experiments.

In US Patent 6,057,134, Ambion describes a method to perform RT-PCRTM which allows an invariant transcript of any relative abundance such as an 18S, 28S, or 5S ribosomal RNA, actin, or glyceraldehyde 3-P phosphate dehydrogenase RNA to be used as a control for any other transcript. This allows two targets of vastly different abundance to be quantified simultaneously in a multiplex RT-PCRTM reaction. Ambion uses blocked primers, or
15 CompetimersTM, that compete with the unmodified primers for binding to a DNA template but cannot be used as primers for extension by a DNA polymerase. Thus, at each extension step in PCRTM, a percentage of template is unavailable for amplification. By increasing the ratio of CompetimersTM to primers in a PCRTM reaction, the amplification efficiency of an amplicon can be reduced so
20 that the linear phase of accumulation of PCRTM product matches that of a less abundant target in multiplex PCRTM.

For a control to be usable for microarray hybridization, the intensity of the signal should be in the same dynamic range as the cDNA under evaluation. rRNA-derived cDNA has never previously proved useful as a control for microarrays
25 probably because it is thousands of times too abundant compared to specific cDNA.

OBJECTS OF THE INVENTION

30 An object of the present invention is therefore to provide an improved method for providing an internal standard for normalizing the relative intensities of signals in hybridization arrays, an improved method for normalizing *per se* and a method of hybridizing making use of the improved normalization.

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SUMMARY OF THE INVENTION

More specifically, in accordance with the present invention, there is provided an improved method for providing an internal standard for normalizing the relative intensities of signals in hybridization arrays that is based on the use of ribosomal RNA (rRNA) as this internal standard. Ribosomal RNA has been
5 found to be particularly suitable for this purpose because its abundance, in terms of percentage of total RNA, does not vary through the cell cycle or with a particular treatment.

The method of the present invention may be summarized as follows. On a
10 given DNA microarray, for example, an oligonucleotide specifically recognizing a sequence contained in ribosomal RNA is spotted along with the other DNA probes used to analyze gene expression, as is usual with this technique. The spots therefore essentially consist of capture probes. Ribosomal RNA, being of relatively invariant quantity in terms of percentage relative to total RNA
15 provides a stable quantitative control to evaluate the quantity of other types of RNA. However, since it is also found in massive amounts relative to other RNAs, its level of detection by the technique must be toned down while remaining accurate. To that end, an experimentally-defined quantity of oligonucleotides carrying the same sequence as that of the oligonucleotide
20 capture probe found on a spot of the microarray is added to the hybridization mixture so that the excess signal coming from the labelled rRNA (or from the cDNA generated from the rRNA, if cDNA hybridization is the method selected) is competed out and the signal detected for it is reduced to a range compatible with that of the signal for the other labelled RNAs.

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Specifically, the present invention provides a novel method for providing an internal standard for normalizing the relative intensities of signals on a hybridization array, comprising:

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adding a known quantity of an unlabelled ribosomal nucleic acid competitor probe into a hybridization buffer suitable for the array experiment, the competitor probe characterized in that it has the same sequence as at least portion of a capture probe present in the array for immobilizing ribosomal nucleic acids thereon; and
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5 allowing the competitor probe to compete with a ribosomal capture probe for hybridization to a suitably labelled rRNA-derived cDNA of a cDNA sample, such that a hybridization signal of labelled rRNA-derived cDNA is decreased to a suitable signal dynamic range of detection and the rRNA-derived cDNA of the sample becomes a suitable internal standard for the hybridization array.

10 The method of the present invention may further include:

15 determinating the quantity of hybridized rRNA-derived cDNA;
 and

 comparing the quantity of hybridized rRNA-derived cDNA
15 against standard curves to determine the quantity of cDNA in
 said sample.

20 The present invention further provides a normalization method, wherein the above steps for obtaining an internal standard are reproduced for a test sample using a first label, and for a suitably-labelled reference sample using a second label, and the quantity of hybridized rRNA-derived cDNA originating from the test sample is compared to the quantity of rRNA-derived cDNA originating from the reference sample hybridizing to the same capture probe to provide a normalization factor.

25 The present invention further provides a hybridization array, wherein the above steps for normalizing are reiterated and the normalization factor is used to correct a hybridization signal provided by the binding of a target cDNA of the test sample labelled with the first label to a capture probe specific to said target, which correction makes said hybridization signal directly comparable to a hybridization signal provided by the binding of the same target of the reference sample labelled with the second label to the same capture probe specific to said target.

35 In a preferred embodiment, the rRNA competitor probe is present in a concentration that is about 5 to about 100 times that of the capture probe.

The rRNA-derived cDNA may be labelled by any suitable means, such as by 3' addition of phosphate, or labelling with cyanines, biotin, digoxigenin, fluorescein, a dideoxynucleotide, an amine, a thiol, an azo (N₃) group or fluorine, or any other form of label.

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An array comprising a plurality of spotted cDNA capture probes for binding ribosomal nucleics, alone or in combination with the competitor ribosomal probe in a separate component are further objects of this invention. The method of the present invention is suitable for use in high-throughput screening experiments.

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It may be used for any type of array experiment, including but not limited to the identification of sequences found in the open reading frame of genes coding for transcription factors, such as c-Rel, E2F-1, Egr-1, ER, NFκB, p53, Sp1 and YY1. ✓

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Other objects, advantages and features of the present invention will become more apparent upon reading of the following non restrictive description of preferred embodiments thereof, given by way of example only with reference to the accompanying drawings.

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BRIEF DESCRIPTION OF THE DRAWINGS

In the appended drawings:

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Figure 1: A summary view of the described technology. Any given pool of total cellular RNA is usually composed of 80% ribosomal RNA (rRNA) and 20% messenger RNA (mRNA) and small nuclear RNAs. mRNA (except for the histone genes) is polyadenylated while rRNA never is. Making cDNA from both types of RNA by reverse transcription is possible if using a poly dT primer for mRNA (producing mRNA-derived cDNA, shown by solid arrows) and a specific primer for rRNA (producing rRNA-derived cDNA, shown by dashed arrows). Analysis of mRNA by microarray using the constant rRNA as a standard is made difficult by the relative overabundance of rRNA relative to mRNA; this problem is circumvented by adding to the hybridization mix a rRNA competitor probe which has the same sequence as the microarray's rRNA-cDNA capture

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probe (both shown as lines marked with an "r"). By sequestering the excess rRNA-derived cDNA, the competitor probe brings down the level of hybridizable and hybridized rRNA-derived cDNA to usable levels.

- 5 **Figure 2:** Human ribosomal DNA complete repeating unit (GB accession number #U13360). ETS: externally transcribed spacer. ITS: internally transcribed spacer. IGS: intergenic spacer. The position of a few rRNA probes is shown.
- 10 **Figure 3:** Illustration of spotted DNA capture probes on the slide. The slide used for the described experiment carries 12 probe blocks, identified 1 to 12. In each block there are 7 rows and 16 columns of spots. Each DNA capture probe was spotted in duplicate in an adjacent column (i.e., all odd columns correspond to a duplicate column) so there are 8 different DNA probes in a
- 15 column. There are a total of 1344 spots on the slide, corresponding to duplicates of 463 different DNA capture probes and 209 negative controls (no DNA probe).
- 20 **Figure 4:** Cohybridization of labelled cDNA from Jurkat (reference sample: Cy3-green) and Jurkat-TPA (test sample: Cy5-red). Ratio images exported from GenePix Pro 3.0 (Axon Instruments Inc.) as JPEG (or TIFF) files are 24-bit RGB color.
- 25 **Figure 5:** Cohybridization of labelled cDNA from Jurkat (Cy3-green) and Jurkat-TPA (Cy5-red). Five (5) ng of rRNA competitor probe 2 was added to the hybridization mix to compete for the hybridization of the rRNA-derived cDNA to the attached rRNA cDNA capture probe 2. Ratio images exported from GenePix Pro 3.0 (Axon Instruments Inc.) as JPEG (or TIFF) files are 24-bit
- 30 RGB color.
- 35 **Figure 6:** Cohybridization of labelled cDNA from Jurkat (Cy3-green) and Jurkat-TPA (Cy5-red). Fifty (50) ng of rRNA competitor probe 2 was added to the hybridization mix to compete for the hybridization of the rRNA-derived cDNA to the attached rRNA cDNA capture probe 2 (which has the same sequence as rRNA competitor probe 2). Ratio images exported from GenePix Pro 3.0 (Axon Instruments Inc.) as JPEG (or TIFF) files are 24-bit RGB color.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENT

GLOSSARY

5 In order to provide a clear and consistent understanding of terms used in the present description, a number of definitions are herein provided.

Array : In the context of this invention, an array is a set of different spotted DNA consisting of capture probes for target nucleic acids. Such an array is
10 described in US Patent No. 5,700,637.

Complementary DNA (cDNA) : DNA that has been synthesized from RNA by the effect of the enzyme reverse transcriptase, converting RNA bases into their complements (A to T, U to A, G to C, C to G).
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Cy3, Cy5 : Non-radioactive fluorescent dyes from Amersham Pharmacia Biotech that are widely used for labeling DNA in microarray experiments.

Feature : A feature is a spot (typically of DNA) on a slide. The collection of such
20 features is called a microarray.

Hybridization : The process of joining two complementary strands of DNA, or one strand each of DNA and RNA, to form a double-stranded molecule.

25 **Messenger RNA (mRNA)** : RNA that is used to direct the protein synthesis that is part of gene expression. It represents but a small fraction of the total RNA found in a cell.

mRNA-derived cDNA : cDNA synthesized from a mRNA template using
30 reverse transcriptase and a mRNA-specific primer.

Microarray-sequestered DNA or DNA capture probe: DNA (single-stranded or double-stranded) that are anchored onto the solid surface of a microarray. (See fuller description of microarrays immediately following this Glossary.)
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Oligonucleotide : A short strand of single-stranded DNA, typically composed of up to 50 bases.

Pixel Intensity : The raw intensity of a pixel on a GenePix (Axon Instrument Inc.) single-wavelength or ratio image, falling in a range from 0 to 65535.

5 **PMT** : Photomultiplier tubes in scanners used to analyze array images. These array images are the end products of comparative hybridization experiments.

Ratio Image : The ratio image is an RGB (Red-Green-Blue) overlay image. In this image, wavelength #1 (635 nm) is mapped to the green channel of the RGB image, and wavelength #2 (532 nm) is mapped to the red channel.
10 Superimposing these two images onto each other results in a third, composite image, whose color is a blend of the red and green signals.

Ratio of medians : The ratio of medians is the ratio of the background subtracted median pixel intensity at the second wavelength to the background subtracted median pixel intensity at the first wavelength.
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Reference cDNA : this cDNA originates from a reference sample that is used for comparison with another one, called test cDNA obtained from a test sample. The reference cDNA serves as a control against which test cDNAs may be compared to quantify changes in the level of expression of any mRNA found in the test sample. Typically, the reference cDNA is labelled with Cy3-dCTP (green fluorescent label) when a fluorescent label is used.
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RGB: Red-Green-Blue color.
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Ribosomal RNA (rRNA): structural RNA found in the ribosomes. It is the most abundant form of RNA in the cell and does not vary significantly.

rRNA-cDNA probe: a probe which is designed to hybridize to the rRNA-derived cDNA found in the hybridization mixture. This probe may be the capture probe, which may have the same sequence as the rRNA competitor probe (see below) so as to compete with it for the target rRNA-derived cDNA.
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rRNA competitor probe: a DNA oligonucleotide with the same sequence as part of a ribosomal RNA-cDNA sequence and capable of competing with the microarray capture probe for hybridization with a rRNA-derived cDNA. This oligonucleotide has the role of competing for the limited space available on the rRNA cDNA capture probe bound to the microarray, thus reducing the quantity
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of rRNA-derived cDNA which can be retained on the microarray and thus allowing the use of rRNA-derived cDNA as an « internal standard ».

5 **rRNA-derived cDNA:** cDNA synthesized from a rRNA template using reverse transcriptase and a rRNA-specific primer.

10 **Saturation :** Saturation refers to the overloading of the photodetection circuitry. Saturation can be reduced by reducing the amount of light that is reaching the PMTs, which is done by reducing the amount of incident laser light. In practice, this is accomplished by reducing the voltage of the PMT, which reduces its gain. Saturating pixels in GenePix 1.0 are shown as white pixels in the raw wavelength images.

15 **Spotted DNA :** Known DNA capture probe that is spotted onto a microarray slide and used to identify the nucleic acids present in unknown samples (test and reference). The spotted DNA could be oligonucleotide or cDNA.

20 **Test cDNA :** cDNA from a cell sample that is to be tested, in comparison with a reference sample. Typically, the test cDNA is labelled with Cy5-dCTP (red fluorescent label) when a fluorescent label is used.

25 Microarrays are made from a collection of purified DNAs. A drop of each type of DNA in solution is placed onto a specially-prepared glass microscope slide by an arraying machine. The arraying machine can quickly produce a regular grid of thousands of spots in a square about 2 cm on a side, small enough to fit under a standard slide cover slip. The DNA in the spots is bound to the glass to keep it from washing off during the hybridization reaction. The choice of DNA to be used within the spots on a microarray's surface determines which genes can be detected in a comparative hybridization assay. These DNA probes could be synthetic oligonucleotides or PCR amplified DNA (hence the terms "oligo
30 microarray" and "cDNA microarray").

The invention relates to rRNA used as an internal standard for the normalization of the fluorescence intensities in microarray analysis experiments. This can provide an estimate of relative abundance of multiple mRNAs and allow direct comparison between two RNA samples.

Use of rRNA for normalization provides a sound method of identifying differentially expressed genes between two samples because its percentage of abundance in total RNA does not vary through the cell cycle or with a particular treatment.

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In order to detect the difference in gene expression between two samples on a single microarray slide, the RNA should be reverse transcribed to cDNA and labelled with two different fluorophores prior to cohybridizing both samples to the same slide and same spots simultaneously. There are several techniques that allow labeling of cDNA. Direct labeling is done by the incorporation of a fluorescent nucleotide such as, for example, Cy3-dCTP (green) or Cy5-dCTP (red) (from Amersham-Pharmacia Biotech), during the reverse transcription reaction. Other protocols may be used for labeling the cDNA following the reverse transcription reaction (indirect labeling). Alternatively, the cDNA can be used for RNA amplification involving T7 polymerase. This method relies on attaching a T7 promoter sequence to the reverse transcriptase primer used for synthesis of the first cDNA strand. After second strand cDNA synthesis, one can generate amplified RNA (aRNA) using T7 RNA polymerase and the double-stranded cDNA molecules as targets for the linear amplification. Those targets can then be labelled directly or indirectly.

In the present invention, the reverse transcriptase reaction for the cDNA labeling step involves the use of two kinds of reverse transcriptase primers in the same reaction: an oligo-dT and specific primers for rRNA (5.8S, 18S or 28S rRNA). One set of RNA to be reverse transcribed is all the polyA+ mRNA that is present in the RNA sample, the other set is the rRNA. Both sets are labelled in the same sample with the same label. Random short primer like random hexamers or sets of specific primers could also be used as alternative methods to reverse transcribe all the polyA+ mRNA.

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In a typical experiment, the reference cDNA is labelled with Cy3 and the test cDNA is prepared in the presence of Cy5. Both of these cDNA populations are hybridized to the same spotted DNA capture probes on the microscope slide. After the hybridization and washing steps, the slide is scanned at the appropriate wavelengths and an image is generated for each wavelength. In the derived ratio image, a red spot indicates that the test cDNA for this feature is more abundant than the reference cDNA which means that the test cDNA is being expressed at a level higher than the reference cDNA; a yellow spot

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means that there is no change in the expression level between the two populations of test and reference cDNA. In order to measure changes in gene expression numerically, image analysis software like GenePix 1.0 (Axon Instruments, Inc.) extracts the intensity of a given feature (spot) from an image and performs a number of computations on the raw data. In this kind of comparative analysis, normalization is essential to compensate for variations in RNA isolation techniques, initial quantification errors, tube to tube variation in reverse transcriptase reactions and other experimental variations. That is where the present invention intervenes: normalization is possible upon correcting the green intensity and the red intensity of the spot having the internal standard capture probe to achieve a ratio of 1. This normalization therefore leads to the obtention of a correction factor that is applied to the intensities of signals specific to each reference and test samples.

The end product of a comparative hybridization experiment is a scanned array image. Saturated pixels appear when there are more photons detected than can be processed by the photomultiplier tubes (PMT) of the scanner. This occurs when the amount of hybridized target per shot is too high. Saturated pixels cannot be used for proper measurement of the signal intensity. PMT should then be set to avoid the detection of saturated pixels. As a consequence, this reduces the signal intensity of all other spots and low levels of cDNA will not be detected.

In the present invention, the hybridization step is performed with specific amounts of free rRNA-derived cDNA (competitor probe) added into the hybridization buffer so as to set up a competition for ribosomal cDNA of the test cDNA and of the reference cDNA (if the latter is part of the experiment) with the capture probe. For efficient competition, the competition probe should be nearly identical to the capture probe or have a high level of overlapping sequences therewith. The hybridization efficiency of the rRNA-derived cDNA with the capture probe can be predictably and reproducibly altered. Reducing the hybridization of these internal and abundant targets in microarray experiments has the effect of generating a signal intensity in the same dynamic range of detection as the less abundant targets in microarrays.

The competition is important because the control must be detected at a level similar to the test transcript. If one target is present at a significantly higher concentration than the other, the PMT (laser voltage) has to be reduced to

avoid a saturated signal, with the consequence of reducing all the other signals. The ability to obtain quantitative information for low abundant mRNA will then be lost.

- 5 With the applicants' invention, the normalization factor is computed using the ratio of intensity obtained between the signal detected for the test cDNA and that of the reference cDNA. This ratio should be 1.0. For example, if the ratio is 0.8, a normalization factor of 1.25 would have to be calculated ($1/0.8$). The analyzed data is then corrected using this factor. If the normalization factor is
10 greater than 2 (or less than 0.5) the slide is usually rescanned with other PMT voltage to ensure maximum data integrity.

RESULTS

- 15 The applicants used the products and protocols that are described herein, which results in proper normalization.

Figure 1 illustrates how a given sample (reference or test) is labelled and hybridized to capture probes (a plurality of specific cDNA probed spots and one
20 internal standard probe spot). The labelled ribosomal cDNA is mixed with a competitor probe that is here identical to the capture probe.

Figure 2 illustrates the organization of the rDNA locus. The microarray was made from a collection of synthetic DNA oligonucleotides as DNA probes.

25 Figure 3 illustrates the positions of spotted DNA capture probes on the slide. In order to use the cDNA made from rRNA for normalisation, a DNA capture probe having a sequence that is complementary to the rRNA-derived cDNA has also been spotted on the array slide.

30 Table 1 shows the sequences of two DNA probes designed for that purpose. 3D-Link Activated slides from Surmodics Inc. were used according to the supplier's protocol for the covalent attachment of the 5' amino modified oligonucleotides and prehybridization treatment of the slides. On the DNA
35 microarray used here, each spot contains approximately 0.15ng of bound DNA probe.

The cDNA for microarray analysis was prepared from RNA templates by incorporation of fluorescent-labelled deoxyribonucleotides during first strand cDNA synthesis. 10µg of total RNA extract from Jurkat and Jurkat-TPA cell lines (Geneka Biotechnology) was used. Priming of cDNA synthesis was performed using 2µg of oligo (dT). For each labeling reaction, 50 ng of 18S primer were included to allow reverse transcription of the 18S rRNA. Table 1 shows the sequences of the 18S reverse transcriptase primer. In this experiment, labelled reference cDNA from Jurkat total RNA was prepared using Cy3-dCTP while Jurkat-TPA total RNA was reverse transcribed and labelled using Cy5-dCTP (Amersham Pharmacia Biotech) to produce labelled test cDNA. Reverse transcriptase reactions were performed using the Superscript II reverse transcriptase (LifeTechnologies) enzyme according to the supplier's protocol.

For the hybridization and washing steps the following conditions were used (optimized conditions for 3D-Link Activated slides, Surmodics Inc.). Labelled cDNAs were cohybridized in 5x SSC-0.1% SDS buffer for 16 hours at 45°C. Washing was performed by incubating slides two times 15 minutes in 2x SSC-0.1% SDS at 45°C, one time 5 minutes in 0.2x SSC at room temperature and one time 5 minutes in 0.1x SSC at room temperature. Slides were dried by low speed centrifugation.

The test and reference cDNAs were analyzed through hybridization with the microarray-sequestered cDNA. In this type of experiment, if the test or reference cDNA contains a sequence that is complementary to the DNA on a given spot, that cDNA will hybridize to the spot, where it will be detectable by virtue of its fluorescence.

Figure 4 shows a ratio image of a typical cohybridized cDNA with no internal standard according to the invention. The target cDNAs and the results are listed in Table 2 (see right column). Figures 5 and 6 show counterparts of arrays of Figure 4 but with 5 ng and 50 ng of ribosomal competitor probe, respectively, in accordance with this invention. The results are listed in Table 2, in the middle and left columns, respectively.

Saturated spots were observed for the two rRNA cDNA probes (DNA probe 1 and probe 2). The GenePix 3.0 software (Axon Instruments Inc.) was used to extract the intensity of each feature (hybridized spot) from the image. Table 2

shows the mean value of pixel intensity for each spot. To analyse feature intensity and calculate a ratio, the local background should be subtracted from the median value of the pixel. The method used by GenePix Pro 3.0 for determining the background intensity is a local background subtraction technique. A different background is therefore computed for each individual feature-indicator and the median value of the background pixel intensities are reported (Table 2).

The end product of a comparative hybridization experiment is a scanned array image. Saturated pixels appear when there are more photons detected than the photomultiplier tubes (PMT) of the scanner can process. This occurs when the amount of hybridized cDNA to the spot is too high. Saturated pixels cannot be used for proper measurement of the signal intensity. PMT should then be set to avoid the detection of saturated pixels. As a consequence, this reduces the signal intensity of all other spots, and lower levels of cDNA will not be detected.

Because of the high abundance of the rRNA-derived cDNA relatively to the mRNA-derived cDNA, it is important to reduce its hybridization to the microarray-sequestered DNA. In this invention, the applicants compete the hybridization of the rRNA-derived cDNA to the microarray DNA capture probe by adding a defined amount of rRNA competitor probe in the hybridization buffer, said probe carrying the same sequence as the microarray-bound probe. Five (5) to 100 molar excess of competitor probe relative to the quantity of microarray DNA capture probe is enough to obtain a rRNA-derived cDNA signal intensity in the same dynamic range of detection as the other cDNAs (i.e., test and/or reference mRNA-derived cDNA), which are otherwise present in much lesser quantities in the reaction buffer. The amount of molar excess to be used is essentially a function of the amount of the total RNA used for the assay (for example : 0.2 to 20 µg).

In short, because of their relatively invariant expression across tissues and treatments, 18S and 28S RNA are ideal internal controls for quantitative RNA analysis by microarrays. The current invention describes how to use these rRNAs to that end by compensating, thanks to competition with specific oligos, for their overabundance relative to the mRNA of test and reference cell samples.

The overall exhaustive results of comparison of test and reference cDNAs, normalized in accordance with the method and principles of the present invention, are provided in appendix 1.

- 5 Although the present invention has been described hereinabove by way of preferred embodiments thereof, it can be modified, without departing from the spirit and nature of the subject invention, as defined in the appended claims.

Name	DNA sequences	Positions relative to 5' 18S sequence	Block	Spotted position Column	Row
RT primer	CTTATGACCCGCACTTACTCG	5'-1667-1647-3'	-	-	-
DNA probe 1	CCCGAGCCGCTGGATACCGCAGCTAGGAATAATGGAATA	5'-833-872-3'	8	1	5
			8	2	5
			1	11	6
			1	12	6
DNA probe 2	TCTCGATTCCGTTGGTGGTGCATGGCCGTTCTTAGTT	5'-1308-1647-3'	10	1	5
			10	2	5
			3	11	6
			3	12	6

Table 1

Block	Column	Row	Gene Name	Probe name	Hybridization with 50 ug of probe 2 as competitor		Hybridization with 5 ug of probe 2 as competitor		Hybridization without competitor	
					Ratio of median value	F535 median	Ratio of median value	F535 median	Ratio of median value	F535 median
					Not normalized	18 S	Not normalized	18 S	Not normalized	18 S
8	1	5	18S	probe 1	1.04	27876	0.73	5617	1.01	55181
1	11	6	18S	probe 1	1.00	53217	0.77	30642	1.01	55181
1	12	6	18S	probe 1	1.00	65349	0.77	30642	1.01	55181
6	2	5	18S	probe 2	0.85	21986	0.68	29798	1.01	65167
10	1	5	18S	probe 2	0.83	73	0.79	4808	1.01	65167
10	2	5	18S	probe 2	1.27	31	1.19	1446	1.01	65250
3	11	6	18S	probe 2	1.00	83	1.24	1437	1.01	65250
5	7	6	Beta actin	actin 1	0.78	1159	0.78	2870	1.01	65157
5	8	6	Beta actin	actin 1	0.88	977	0.81	2778	1.01	65157
10	3	1	Beta actin	actin 1	0.87	1674	0.74	2813	1.01	65168
10	4	1	Beta actin	actin 1	0.89	1880	0.86	2142	1.01	65168
4	3	1	Beta actin	actin 1	0.63	2010	0.79	866	1.01	42650
11	14	5	Beta actin	actin 1	0.85	1607	0.85	1246	1.01	42650
11	13	5	Beta actin	actin 1	0.81	1760	0.82	4081	1.01	42650
4	4	1	Beta actin	actin 1	0.86	1833	0.84	4163	1.01	42650
6	1	1	Beta actin	actin 1	0.88	3519	0.82	5178	1.01	42650
3	2	1	Beta actin	actin 1	0.88	278	0.86	861	1.01	42650
3	3	1	Beta actin	actin 1	0.88	1687	0.86	3107	1.01	42650
3	4	1	Beta actin	actin 1	0.88	3013	0.86	3255	1.01	42650
6	2	1	Beta actin	actin 2	0.75	1641	0.75	3905	1.01	42650
4	1	1	Beta actin	actin 2	0.73	1651	0.72	3905	1.01	42650
4	1	1	Beta actin	actin 2	0.91	419	0.86	102	1.01	42650
3	1	1	Beta actin	actin 2	0.87	530	0.87	2931	1.01	42650
5	1	1	Beta actin	actin 2	0.77	323	0.80	1924	1.01	42650
5	2	1	Beta actin	actin 2	0.75	2986	0.83	8491	1.01	42650
3	7	6	Beta actin	actin 3	1.41	1336	1.38	5556	1.01	42650
3	8	6	Beta actin	actin 3	1.26	1744	1.46	5366	1.01	42650
11	2	1	Beta actin	actin 3	1.51	1697	1.73	1674	1.01	42650
11	1	1	Beta actin	actin 3	1.22	572	1.31	2150	1.01	42650
12	2	1	Beta actin	actin 3	1.13	534	1.28	4607	1.01	42650
10	2	1	Beta actin	actin 3	1.11	651	1.18	4478	1.01	42650
9	2	1	Beta actin	actin 3	1.23	947	1.37	1003	1.01	42650
10	1	1	Beta actin	actin 3	0.92	560	1.05	7356	1.01	42650
8	2	1	Beta actin	actin 3	1.28	514	1.26	5489	1.01	42650
8	1	1	Beta actin	actin 3	1.36	991	1.69	1937	1.01	42650
9	13	5	Beta actin	actin 3	1.25	1370	1.67	1996	1.01	42650
9	14	5	Beta actin	actin 3	1.43	1339	1.70	4283	1.01	42650
2	1	1	Beta actin	actin 3	0.76	1946	1.60	2481	1.01	42650
2	2	1	Beta actin	actin 3	0.76	1800	1.18	988	1.01	42650
4	2	1	Beta actin	actin 3	0.69	361	0.98	905	1.01	42650
4	2	1	Beta actin	actin 3	0.69	197	0.82	777	1.01	42650
9	14	4	9S8 splicing	L2253_B	2.30	487	2.86	3379	1.01	42650
4	4	4	A-Myo	X13294_B	1.33	332	3.47	62	1.01	42650
4	4	4	ASH1	L6424_A	3.86	1627	1.25	138	1.01	42650
3	5	3	BTEB	O31716_B	4.15	-136	0.79	296	1.01	42650
3	5	3	BTF3 homologue	M60355_A	0.52	54	1.60	173	1.01	42650
2	4	2	CBFA1/OSF2	AF053649_B	0.45	4205	1.33	145	1.01	42650
2	4	2	CDP	M74099_B	0.45	173	1.33	145	1.01	42650
11	10	5	cydin D1	ANL12	1.75	476	3.61	1635	1.01	42650
6	6	6	EN2	L12700_B	2.93	1517	2.15	3462	1.01	42650
8	15	6	GAPDH	S61_	1.37	2104	1.79	184	1.01	42650
2	10	2	GF2IP1	AF036613_B	0.49	106	0.68	45	1.01	42650
5	12	1	ZRP-1	AF003974_A	2.99	4235	3.24	12043	1.01	42650

Table 2

Appendix 1: Signal normalization using 18S RNA as an internal standard. Two microarray analyses were performed independently, each one comparing the expression of many transcription factors in Jurkat cells and in Jurkat cells treated with the phorbol ester TPA. The signals obtained in the latter case were divided by the signals obtained in the former case to get a ratio of induction by TPA in these cells. The signals were normalized using 18S RNA as a standard (see columns 3 and 4). Since 18S RNA is used as a control in both experiments and that the same type of cells were used, presumably giving very similar results, the ratio of the results obtained in each experiment should be nearing 1. That ratio is presented in column 5.

Column 1	Column 2	Column 3	Column 4	Column 5
Gene name	Accession number	Jurkat/Jurkat TPA ratio experiment 1	Jurkat/Jurkat TPA ratio experiment 2	Ratio of experiments 1 and 2
9G8 splicing factor	L22253	0.84	1.00	0.836078512
9G8 splicing factor	L22253	0.77	0.99	0.779340183
A-Myb	X66087	1.32	1.38	0.950679679
A-Myb	X66087	1.34	1.43	0.937305665
A-Myb	X13294	1.12	1.21	0.924150275
A-Myb	X13294	1.12	1.21	0.924083463
ABF-1	AF060154	0.45	0.39	1.166895465
ABF-1	AF060154	0.39	0.38	1.029207795
ABH	NM_006020	0.91	1.05	0.865303363
ABH	NM_006020	0.81	0.98	0.822950019
ABP/ZF	U82613	1.32	1.64	0.804108596
ABP/ZF	U82613	1.25	1.60	0.783304597
AF10	NM_004641	1.24	1.31	0.947593818
AF10	NM_004641	1.23	1.32	0.931357689
AIB3	AF208227	1.33	1.28	1.034779297
AIB3	NM_014071	1.09	1.25	0.870698314
AIB3	NM_014071	1.07	1.36	0.784035932
AIB3	AF208227	1.10	1.40	0.782294079
ALL-1	U04737	1.65	1.88	0.880126672
ALL-1	U04737	1.58	1.88	0.838592996
ALL-1	L04284	0.66	0.79	0.838134698
AML2	Z35278	0.44	0.51	0.858684813
AML2	Z35278	0.42	0.55	0.77112205
AML3	AF001450	1.28	1.32	0.974983445
AML3	AF001450	1.34	1.39	0.966458433
AP-2gamma	U85658	2.57	2.62	0.978390776
AP-2gamma	U85658	2.23	2.59	0.86381938
AP-4	X57435	1.21	1.23	0.984438472

AP-4	X57435	1.17	1.28	0.91144528
AP4	NM_014374	1.39	1.59	0.871879245
AP4	NM_014374	1.32	1.59	0.831996755
APBB1	NM_001164	0.95	0.97	0.984113563
APBB1	NM_001164	0.79	0.99	0.801180869
APC	M74088	1.50	1.31	1.148676257
APC	M74088	1.29	1.46	0.8859936
APECED	AB006682	1.49	1.56	0.957659838
APECED	AB006682	1.38	1.65	0.837168643
APEX	NM_001641	0.88	1.13	0.783250131
APEX	NM_001641	0.84	1.08	0.780343345
APOBEC2	NM_006789	1.15	1.12	1.031439776
APOBEC2	NM_006789	1.04	1.05	0.990111417
APPL	NM_012096	1.32	1.54	0.856820461
APPL	NM_012096	1.31	1.56	0.839878811
AR	NM_000044	1.74	2.04	0.855879355
AR	NM_000044	1.60	2.01	0.796494966
ARNT	M69238	1.25	1.42	0.880056649
ARNT	M69238	1.24	1.42	0.876705905
ARNT	Y18500	0.78	0.96	0.816130578
ASH2L2	AF056717	1.34	1.35	0.994678817
ASH2L2	AF056717	1.38	1.40	0.991252318
ATBF1	NM_006885	0.90	1.01	0.889758762
ATBF1	NM_006885	0.90	1.02	0.879456944
ATF	D90209	1.05	1.01	1.035713928
ATF	D90209	0.97	1.01	0.960323304
ATF-a	X52943	1.54	1.88	0.817277421
ATF-a	X52943	1.51	1.93	0.780957523
ATF1	NM_005171	0.84	0.91	0.927916867
ATF1	NM_005171	0.87	1.02	0.854281302
ATF6	NM_007348	1.29	1.29	1.00327664
ATF6	NM_007348	1.09	1.28	0.856533977
BACH1	NM_001186	1.49	1.31	1.137064444
BACH1	NM_001186	1.45	1.62	0.891057108
BAPX1	NM_001189	2.55	2.33	1.093826453
BAPX1	NM_001189	2.46	2.59	0.946872482
BARX2	NM_003658	1.17	1.27	0.917084438
BARX2	NM_003658	1.14	1.37	0.830998058
BCL2	NM_000633	1.43	1.65	0.866945304
BCL2	NM_000633	1.37	1.70	0.806442848
BCL3	U05822	1.11	1.26	0.877431885
BCL3	M31732	1.17	1.38	0.848343893
BCL3	M31732	1.13	1.37	0.825031918
BCL3	U05822	1.02	1.30	0.790257156
Beta-actin	X00351	1.02	1.19	0.855958172
Beta-actin	X00351	1.02	1.21	0.843968769
Beta-actin	X00351	1.01	1.21	0.837209294
Beta-actin	X00351	1.00	1.19	0.836410947
beta-catenin	X89593	2.01	2.06	0.977986591
beta-catenin	X89593	1.99	2.11	0.942592932
BF-2	X74143	1.28	1.38	0.931388014
BF-2	X74143	1.22	1.37	0.894927517

BFP/ZNF179	AB026054	1.33	1.32	1.005754548
BFP/ZNF179	AB026054	1.36	1.37	0.993222418
BIRC4	NM_001167	1.51	1.44	1.054435009
BIRC4	NM_001167	1.40	1.50	0.932289706
BMZF3	NM_005773	0.92	1.08	0.850837495
BMZF3	NM_005773	0.90	1.13	0.798215326
brahma	X72889	5.90	5.49	1.074544412
brahma	X72889	5.14	5.97	0.86166573
BRCA2	NM_000059	1.45	1.75	0.824507422
BRCA2	NM_000059	1.39	1.74	0.798236353
Brn-3B	U06233	1.48	1.37	1.078166711
Brn-3B	U06233	1.47	1.50	0.974841891
Brn-4	X82324	1.57	1.06	1.486851514
Brn-4	X82324	1.29	1.07	1.198217087
BRS3	NM_001727	2.71	2.75	0.983814035
BRS3	NM_001727	2.36	2.77	0.851828571
BTEB	D31716	4.86	4.21	1.153934489
BTEB	D31716	4.30	4.32	0.995197771
BTEB2	D14520	1.25	1.27	0.978590601
BTEB2	D14520	1.30	1.39	0.933625786
BTF3	NM_001207	1.05	1.10	0.955111894
BTF3	NM_001207	0.99	1.08	0.913787418
BTF3a	M90352	2.83	2.32	1.219855319
BTF3a	M90352	2.70	2.39	1.130461687
BTF3L1	NM_001208	1.22	1.07	1.137813523
BTF3L1	NM_001208	1.16	1.05	1.102860167
BTF3L3	M90356	1.44	1.37	1.049188317
BTF3L3	M90356	1.24	1.34	0.927268611
bZip protein B-ATF	U15460	1.07	1.14	0.9426678
bZip protein B-ATF	U15460	0.97	1.08	0.901877866
c-Ets-1	X14798	1.09	1.25	0.873492353
c-Ets-1	X14798	1.10	1.32	0.830363686
c-maf	AF055376	5.74	4.79	1.19705637
c-maf	AF055376	4.91	5.10	0.962031195
c-Rel	M11595	1.33	1.41	0.946493027
c-Rel	X75042	1.32	1.46	0.902036285
c-Rel	M11595	1.27	1.42	0.889929469
c-Rel	X75042	1.14	1.47	0.777782886
C2H2 ZNF	AF033199	1.07	1.14	0.938338671
C2H2 ZNF	AF033199	0.99	1.16	0.852890579
C2H2-type ZNF	U95991	1.19	1.01	1.173282928
C2H2-type ZNF	U95991	0.98	1.04	0.942590144
C2ORF3	NM_003203	1.46	1.22	1.196699322
C2ORF3	NM_003203	1.01	0.93	1.093811577
CBF (5)	M37197	4.06	4.25	0.956014195
CBF (5)	M37197	3.60	4.09	0.88090602
CBF1	AF098297	1.61	1.63	0.991664197
CBF1	AF098297	1.38	1.78	0.772546908
CBFA1	L40992	1.30	1.45	0.898057655
CBFA1	L40992	1.26	1.46	0.865127809
CBFA1/OSF2	AF053949	1.22	1.28	0.951727989
CBFA1/OSF2	AF053949	1.22	1.33	0.92146037

CBFA2T1	NM_004349	1.49	1.65	0.901008111
CBFA2T1	NM_004349	1.24	1.59	0.780002118
CBFB	L20298	2.33	2.74	0.851333501
CBFB	L20298	2.36	2.91	0.8088749
CDP	M74099	1.39	1.61	0.85914075
CDP	M74099	1.27	1.64	0.77621359
CEBPB	NM_005194	1.24	1.47	0.846246886
CEBPB	NM_005194	1.26	1.49	0.846246188
CEBPD	NM_005195	0.83	1.00	0.829917576
CEBPD	NM_005195	0.84	1.03	0.822579365
CEBPE	U48866	1.91	2.01	0.948532903
CEBPE	U48866	2.06	2.38	0.86669978
CEZANNE	NM_020205	2.88	2.96	0.974633442
CEZANNE	NM_020205	2.65	2.83	0.935357017
CHD1	NM_001270	1.62	1.59	1.014951939
CHD1	NM_001270	1.43	1.59	0.898362477
CHD4	NM_001273	1.54	1.70	0.909055986
CHD4	NM_001273	1.49	1.72	0.862018232
CHFR	NM_018223	4.35	4.43	0.982194772
CHFR	NM_018223	3.92	4.36	0.899117503
CHN1	NM_001822	1.42	1.53	0.927629676
CHN1	NM_001822	1.37	1.49	0.923095091
CIS4	NM_004232	1.67	1.79	0.935688257
CIS4	NM_004232	1.82	2.13	0.851569476
CITED1	NM_004143	1.10	1.30	0.850853943
CITED1	NM_004143	1.17	1.39	0.844249881
CNBP	M28372	0.67	0.54	1.233592517
CNBP	M28372	0.62	0.54	1.163359863
coactivator EBV nuclear protein 2	U22055	0.82	0.94	0.869546763
coactivator EBV nuclear protein 2	U22055	0.81	1.00	0.810099254
COPEB	NM_001300	1.14	1.29	0.885046712
COPEB	NM_001300	1.12	1.34	0.833843243
COPS5	NM_006837	2.46	2.14	1.148421053
COPS5	NM_006837	2.48	2.32	1.071355007
CP2	U01965	1.01	1.23	0.82004865
CP2	U01965	1.00	1.30	0.771414141
CR53	AF017433	1.33	1.33	0.997732351
CR53	AF017433	1.29	1.39	0.925956448
CRE-BP1	J05623	1.13	1.38	0.819277436
CRE-BP1	J05623	1.02	1.26	0.815059942
CREB	M27691	0.92	1.09	0.842697518
CREB	M27691	0.85	1.06	0.7964146
CREBBP	NM_004380	1.09	1.25	0.872661186
CREBBP	NM_004380	1.12	1.30	0.86705145
CREBPA	NM_004904	1.26	1.30	0.971711147
CREBPA	NM_004904	1.10	1.24	0.887551154
CROC4	NM_006365	1.15	1.25	0.926055212
CROC4	NM_006365	1.16	1.38	0.842320854
CRSP70	NM_004831	0.91	1.06	0.854668195
CRSP70	NM_004831	0.92	1.15	0.803384327
CRSP9	NM_004270	1.37	1.49	0.919973517
CRSP9	NM_004270	1.40	1.53	0.919262135

EKLF	U65404	0.96	1.03	0.935317019
ELF1	M82882	1.76	1.83	0.964878433
ELF1	M82882	1.62	1.76	0.921751518
ELF4	NM_001421	1.45	1.41	1.027947336
ELF4	NM_001421	1.36	1.37	0.991044834
ELK3	NM_005230	1.28	1.57	0.815739725
ELK3	NM_005230	1.33	1.68	0.790796088
ELL	NM_006532	0.95	1.16	0.822566492
ELL	NM_006532	0.95	1.16	0.819455294
elongation factor 1-	X16869	1.35	1.50	0.8947725
alpha elongation factor 1-	X16869	1.36	1.59	0.853485168
alpha elongation factor SIII	L34587	1.41	1.64	0.861800291
elongation factor SIII	L34587	1.49	1.82	0.820065033
elongation factor-1-	Z21507	0.81	0.99	0.81190776
delta elongation factor-1-	Z21507	0.78	1.00	0.782148893
delta elongation factor-1-	Z21507	0.78	1.00	0.782148893
EN1	L12698	1.36	1.45	0.935865444
EN1	L12698	1.23	1.47	0.836794344
EPAS1	NM_001430	1.18	1.38	0.856844874
EPAS1	NM_001430	1.15	1.46	0.783761416
ERCC2	X52222	5.72	4.80	1.193231705
ERCC2	X52222	5.33	4.73	1.127089247
ERCC3	NM_000122	1.36	1.57	0.863467286
ERCC3	NM_000122	1.30	1.60	0.812147676
ERF-2	X78992	2.14	2.41	0.889330713
ERF-2	X78992	2.26	2.55	0.883602051
ERG	NM_004449	1.62	1.42	1.142428678
ERG	NM_004449	1.49	1.50	0.996969892
ERM	X96375	4.16	4.29	0.969559654
ERM	X96375	3.27	3.55	0.921520209
ERT	AF017307	2.43	2.68	0.90894817
ERT	AF017307	2.51	2.82	0.891141057
ESRRG	NM_001438	0.95	1.13	0.839582135
ESRRG	NM_001438	0.95	1.15	0.821231854
ETR101	NM_004907	2.74	2.75	0.997375352
ETR101	NM_004907	2.49	2.80	0.887790293
Ets transcription factor	AF115403	1.14	1.31	0.87442124
ESE-2b				
Ets transcription factor	AF115403	1.11	1.43	0.77156259
ESE-2b				
Ets-1 gene	AF193068	1.21	1.38	0.874625305
Ets-1 gene	AF193068	1.22	1.40	0.868962372
Ets-like	U30174	1.40	1.23	1.131217765
Ets-like	U30174	1.49	1.35	1.098811633
Ets-like	Z49980	1.61	1.51	1.067048232
Ets-like	Z49980	1.54	1.56	0.991710772
Ets2	M30137	1.75	2.02	0.86945137
Ets2	M30137	1.78	2.11	0.844919404
ETV1	NM_004956	1.13	1.25	0.910122678
ETV1	NM_004956	1.39	1.59	0.871215971
ETV6	U45432	1.38	1.43	0.965065589
ETV6	NM_001987	0.90	1.11	0.811726255
Evi-1	S82592	2.53	2.10	1.208239627

Evi-1	S82592	2.26	2.15	1.055074375
EWSR1	NM_005243	1.01	1.28	0.789906804
EWSR1	NM_005243	1.00	1.28	0.783731221
EZH2	U61145	1.26	1.35	0.932953273
EZH2	U61145	1.27	1.39	0.907474288
FACTP140	NM_007192	1.43	1.48	0.96265369
FACTP140	NM_007192	1.41	1.48	0.954817504
Fas-binding Daxx	protein AF015956	0.90	1.08	0.833884369
Fas-binding Daxx	protein AF015956	0.89	1.09	0.81465638
FBW1A	AF129530	1.31	1.45	0.900471742
FBW1A	AF129530	1.27	1.54	0.829306514
FGD1	U11690	1.33	1.14	1.173441119
FGD1	U11690	1.21	1.23	0.990554056
FGR	NM_005248	1.33	1.58	0.839283541
FGR	NM_005248	1.27	1.60	0.790883893
FHL1	AF110763	1.56	1.77	0.88200997
FHL1	AF110763	1.45	1.76	0.822210318
FKHL7	AF048693	3.42	3.29	1.040543697
FKHL7	AF048693	3.65	3.62	1.006927826
FKHR	AF032885	2.42	2.08	1.161966778
FKHR	AF032885	2.36	2.18	1.082816723
FKHRL1P1	AF032887	1.42	1.54	0.924383924
FKHRL1P1	AF032887	1.46	1.60	0.912174436
FLI_CDNA	AL360183	1.33	1.28	1.036167415
FLI_CDNA	AL360183	1.37	1.37	0.996443864
FLJ10173	NM_018014	1.04	1.04	0.999229429
FLJ10173	NM_018014	1.01	1.01	0.996944727
FLJ10251	NM_018039	1.31	1.43	0.911977997
FLJ10251	NM_018039	1.31	1.46	0.897214657
FLJ10339	NM_018063	1.62	1.87	0.866263178
FLJ10339	NM_018063	1.53	1.86	0.822236451
FLJ10469	NM_018102	0.94	1.09	0.865336872
FLJ10469	NM_018102	0.94	1.09	0.865236102
FLJ10688	AK001550	0.97	1.11	0.881574929
FLJ10688	AK001550	0.95	1.19	0.802514491
FLJ10891	NM_018260	1.22	1.31	0.928051813
FLJ10891	NM_018260	1.15	1.38	0.83802715
FLJ10909	AK001771	2.36	2.38	0.988865574
FLJ10909	AK001771	2.19	2.39	0.917728349
FLJ11015	NM_018300	1.10	1.19	0.922727928
FLJ11015	NM_018300	0.99	1.17	0.845365497
FLJ11137	NM_018337	1.43	1.64	0.875337744
FLJ11137	NM_018337	1.33	1.60	0.831187459
FLJ11340	AK002202	3.74	3.96	0.944037815
FLJ11340	AK002202	3.72	4.02	0.925252175
FLJ11344	AK002206	1.11	1.24	0.900786005
FLJ11344	AK002206	1.22	1.39	0.879708376
FLJ11688	AK021750	1.31	1.41	0.925531252
FLJ11688	AK021750	1.35	1.50	0.901037884
FLJ12606	AK022668	1.11	1.15	0.965529216
FLJ12606	AK022668	0.96	1.10	0.876202729
FLJ12628	AK022690	1.30	1.38	0.938935506

FLJ12628	AK022690	1.25	1.36	0.925116959
FLJ12644	AK000909	0.98	1.09	0.901825689
FLJ12644	AK000909	1.02	1.16	0.874104763
FLJ13479	AK023541	1.12	1.35	0.830838509
FLJ13479	AK023541	1.05	1.27	0.825925564
FLJ20337	NM_017772	1.55	1.60	0.969110576
FLJ20337	NM_017772	1.60	1.66	0.966477023
FLJ20428	AK000435	1.00	1.13	0.88699187
FLJ20428	AK000435	1.03	1.16	0.883146291
FLJ20438	ak000445	2.61	2.97	0.876697181
FLJ20438	ak000445	2.41	2.99	0.807812353
FLJ22332	AK025985	1.37	1.67	0.823105199
FLJ22332	AK025985	1.22	1.52	0.80219778
FLJ22973	AK026626	0.93	1.11	0.841359026
FLJ22973	AK026626	0.94	1.14	0.825377156
FOG2	NM_012082	1.03	1.10	0.930301277
FOG2	NM_012082	1.11	1.24	0.901732208
FOSL2	NM_005253	1.42	1.73	0.818161857
FOSL2	NM_005253	1.40	1.80	0.775807396
FOXD2	NM_004474	1.36	1.49	0.918399567
FOXD2	NM_004474	1.35	1.48	0.912187342
FOXD3	NM_012183	1.67	1.55	1.072311149
FOXD3	NM_012183	1.55	1.63	0.952188792
FOXO3A	NM_001455	0.87	1.10	0.789948212
FOXO3A	NM_001455	0.85	1.09	0.780082817
FRA-1	X16707	1.19	1.22	0.975373174
FRA-1	X16707	1.15	1.25	0.920368654
FREAC1	U13219	1.33	1.44	0.920850002
FREAC1	U13219	1.33	1.47	0.900355759
FREAC10	AF042831	1.37	1.53	0.895510594
FREAC10	AF042831	1.29	1.49	0.862685753
FREAC6	L13203	0.68	0.76	0.894745854
FREAC6	L13203	0.65	0.75	0.865206105
FREAC7	U13225	0.82	0.70	1.159351607
FREAC7	U13225	0.70	0.72	0.971169803
frpHE	AF026692	1.02	1.11	0.917414227
frpHE	AF026692	0.99	1.17	0.852068204
GABPB1	NM_005254	1.74	1.77	0.98532103
GABPB1	NM_005254	1.74	1.82	0.956282084
GADD 153	s40706	1.28	1.47	0.871741003
GADD 153	s40706	1.09	1.42	0.76973024
GAPDH	M33197	0.58	0.61	0.949162972
GAPDH	M33197	0.56	0.59	0.947048611
GCMA	NM_003643	1.20	1.32	0.908325507
GCMA	NM_003643	1.15	1.32	0.870108289
GCN5L1	NM_001487	0.80	0.93	0.856784201
GCN5L1	NM_001487	0.73	0.90	0.820087548
GIOT-1	AB021641	1.29	1.45	0.884372648
GIOT-1	AB021641	1.22	1.50	0.812297818
GIOT-2	NM_016264	0.93	1.07	0.868101488
GIOT-2	NM_016264	0.91	1.10	0.82720992
GIOT-3	NM_016265	0.87	0.97	0.893216374

GIOT-3	NM_016265	0.86	0.97	0.884789805
GIOT-4	NM_016266	1.73	2.15	0.806073313
GIOT-4	NM_016266	1.78	2.27	0.783305867
GLI	X07384	1.34	1.32	1.019171414
GLI	X07384	1.28	1.34	0.958829722
GLI3	M57609	2.18	1.98	1.098071683
GLI3	M57609	1.95	2.06	0.944192578
GPX5	NM_001509	0.94	1.08	0.865100605
GPX5	NM_001509	1.37	1.62	0.847040407
GRLF1	NM_004491	0.79	0.87	0.906709069
GRLF1	NM_004491	0.71	0.80	0.885942625
GTF2B	NM_001514	2.16	2.30	0.937394785
GTF2B	NM_001514	1.95	2.44	0.799673306
GTF2E1	NM_005513	0.76	0.98	0.784112092
GTF2E1	NM_005513	0.74	0.96	0.769972038
GTF2I	NM_001518	2.68	2.80	0.958900201
GTF2I	NM_001518	2.58	2.96	0.869928806
GTF2IP1	AF036613	0.95	0.70	1.359169172
GTF2IP1	AF036613	0.84	0.84	0.988769754
GTF3A	NM_002097	1.49	1.59	0.932083753
GTF3A	NM_002097	1.58	1.70	0.931723008
GTF3C1	NM_001520	2.07	2.25	0.919747554
GTF3C1	NM_001520	1.99	2.34	0.847308047
GTF3C2	NM_001521	1.31	1.28	1.027107124
GTF3C2	NM_001521	1.22	1.29	0.943844023
GTF3C3	NM_012086	1.64	1.64	0.996581398
GTF3C3	NM_012086	1.52	1.67	0.907893639
GTF3C4	NM_012204	1.11	1.28	0.860437792
GTF3C4	NM_012204	1.13	1.33	0.851773956
GTP	AF054183	1.98	2.25	0.880760132
GTP	AF054183	1.86	2.27	0.818437867
H1F3	M60746	1.08	1.27	0.853497342
H1F3	M60746	1.10	1.33	0.824710104
H2AFX	X14850	1.24	1.33	0.934617922
H2AFX	X14850	1.19	1.40	0.849848259
H4	X67081	0.83	0.97	0.859373264
H4	X67081	0.84	1.00	0.842811776
hairless	AF039196	1.39	1.46	0.951801096
hairless	AF039196	1.37	1.53	0.896272718
HAP2	M59079	1.59	1.49	1.062457371
HAP2	M59079	1.33	1.71	0.780793131
HAT1	NM_003642	1.05	0.86	1.223229142
HAT1	NM_003642	1.06	1.04	1.026464835
HB16	M31630	0.97	1.09	0.894291244
HB16	M31630	1.01	1.26	0.797052293
HB9	U07663	2.64	2.61	1.013508831
HB9	U07664	0.92	1.03	0.895489189
HBOA	NM_007067	1.27	1.37	0.929935594
HBOA	NM_007067	1.23	1.35	0.912294782
HCF-2	AF117210	1.43	1.56	0.918694023
HCF-2	AF117210	1.43	1.61	0.888145397
HD-ZNF1	NM_004876	1.11	1.22	0.910541692

HD-ZNF1	NM_004876	1.05	1.19	0.884372648
HDAC1	NM_004964	0.83	0.97	0.851140233
HDAC1	NM_004964	0.82	0.97	0.844737874
HDAC4	NM_006037	2.76	2.40	1.153229661
HDAC4	NM_006037	1.64	1.99	0.825094678
HEB	M83233	0.75	0.89	0.83812814
HEB	M83233	0.73	0.90	0.814864061
HEN1	M96739	1.51	1.61	0.937658625
HEN1	M96739	1.49	1.69	0.883530062
HERP1	AF232238	1.64	1.78	0.918811847
HERP1	AF232238	1.48	1.69	0.873182906
HERP2	AF232239	0.88	0.97	0.913791819
HERP2	AF232239	0.82	1.01	0.814129508
HES4	AB048791	1.12	1.24	0.906421263
HES4	AB048791	1.17	1.31	0.892326717
HGS	NM_004712	1.13	1.22	0.925941974
HGS	NM_004712	1.08	1.23	0.884627755
HIC1	NM_006497	1.01	1.19	0.84718439
HIC1	NM_006497	0.94	1.20	0.789513268
HIVEP1	NM_002114	1.24	1.14	1.08520656
HIVEP1	NM_002114	1.03	1.16	0.893216374
HIVEP2	NM_006734	2.86	2.87	0.99372865
HKE4	NM_006979	1.51	1.70	0.89115193
HKE4	NM_006979	1.35	1.66	0.814320291
HLF	M95585	1.28	1.32	0.971118298
HLF	M95586	1.15	1.26	0.910803018
HMG-1	D63874	1.27	1.25	1.015741343
HMG-1	D63874	1.23	1.22	1.008346304
HMG-2	X62534	1.70	1.82	0.938295788
HMG-2	X62534	1.55	1.76	0.878616998
HMG17	NM_005517	0.99	1.14	0.868604212
HMG17	NM_005517	0.97	1.15	0.841273347
HMG1Y	NM_002131	0.92	1.04	0.886466628
HMG1Y	NM_002131	0.93	1.13	0.824826257
HNF-1A	M57732	1.03	1.19	0.868596298
HNF-1A	M57732	1.07	1.25	0.861349263
HNF-1B	X71346	2.40	2.21	1.087117438
HNF-1B	X71346	2.25	2.18	1.030922798
HNF-3gamma	L12141	1.46	1.53	0.956635501
HNF-3gamma	L12141	1.40	1.54	0.90844598
HNF-4alpha3	U72967	2.92	3.06	0.953909282
HNF-4alpha3	U72967	2.76	3.16	0.871764387
HNF-6alpha	AF035580	1.20	1.00	1.202677165
HNF-6alpha	AF035580	1.02	1.07	0.954515537
HNF3A	NM_004496	1.35	1.39	0.968770391
HNF3A	NM_004496	1.30	1.39	0.934312714
HOX	L11239	1.29	1.55	0.831459424
HOX	L11239	1.22	1.56	0.784287548
HOX11	s38742	0.82	0.97	0.846268344
HOX11	s38742	0.89	1.06	0.840219605
HOX11L2	AJ223798	5.90	5.44	1.08601856
HOX11L2	AJ223798	5.29	5.47	0.967027069

HOXA-9	U81511	2.28	2.06	1.107860869
HOXA-9	U81511	2.06	2.02	1.019494694
HOXA1	S79910	1.47	1.44	1.023612925
HOXA1	S79910	1.22	1.31	0.930731462
HOXA11	AF071164	1.23	1.36	0.902672948
HOXA11	AF071164	1.28	1.48	0.86247018
HOXA13	NM_000522	7.13	5.19	1.375914112
HOXA13	NM_000522	3.90	4.45	0.876388041
HOXA4	U56105	1.20	1.41	0.854164123
HOXA4	U56105	1.19	1.46	0.814779811
HOXA7	NM_006896	1.14	1.20	0.952764133
HOXA7	NM_006896	1.09	1.21	0.899003953
HOXB1	X16666	1.59	1.81	0.877682176
HOXB1	X16666	1.62	2.00	0.80887332
HOXB2	X78978	1.84	1.60	1.145917
HOXB2	X78978	1.64	1.72	0.957991608
HOXB2	X16665	1.39	1.54	0.905368978
HOXB2	X16665	1.42	1.59	0.895429132
HOXB3	X16667	1.92	1.73	1.107588304
HOXB3	X16667	1.87	1.84	1.015740013
HOXB4	AF005652	1.16	1.27	0.911652213
HOXB4	AF005652	1.09	1.24	0.880915725
HOXB5	M92299	1.18	1.38	0.854344138
HOXB5	M92299	1.20	1.49	0.803737757
HOXB7	M16937	0.95	1.22	0.778800068
HOXB7	M16937	0.97	1.24	0.778387715
HOXC10	AF255675	1.16	1.28	0.905053085
HOXC10	X99685	1.12	1.27	0.881270065
HOXC10	AF255675	1.13	1.31	0.858450467
HOXC10	X99685	1.10	1.33	0.82796661
HOXC6	M16938	1.26	1.49	0.844466889
HOXC6	M16938	1.16	1.46	0.800039127
HOXC8	X99681	1.12	1.30	0.860768554
HOXC8	X99681	0.97	1.24	0.783209726
HOXD3	NM_006898	1.51	1.62	0.92856985
HOXD3	NM_006898	1.47	1.61	0.918716026
HOXD4	X04706	1.24	1.40	0.886519344
HOXD4	X67079	1.56	1.78	0.877418885
HOXD4	X67079	1.54	1.86	0.826005297
HOXD4	X04706	1.22	1.52	0.804048475
HPX42B	NM_014468	1.02	1.04	0.980963071
HPX42B	NM_014468	0.91	1.00	0.913143774
hRev	X72631	1.25	1.35	0.929674185
hRev	X72631	1.28	1.42	0.902255362
HS747E2A	NM_015370	1.07	1.12	0.959032318
HS747E2A	NM_015370	1.02	1.17	0.873166624
HSA275986	NM_018403	1.80	1.66	1.081002809
HSA275986	NM_018403	1.61	1.81	0.888060724
HSBP1	AF068754	2.24	2.62	0.853507954
HSBP1	AF068754	2.27	2.83	0.801085361
HSET	D14678	0.47	0.56	0.84140568
HSET	D14678	0.46	0.59	0.779570541

HSF2BP	NM_007031	2.36	2.61	0.904409562
HSF2BP	NM_007031	2.24	2.57	0.86866997
HSGT1	NM_007265	1.14	1.17	0.973056944
HSGT1	NM_007265	1.12	1.27	0.878498082
hSIM2	D85922	2.71	2.85	0.952407887
hSIM2	D85922	2.65	2.91	0.910509622
Hsp90	X07270	0.92	1.11	0.82588322
Hsp90	X15183	2.01	2.48	0.812100632
hTFIIS.h	AJ223473	0.99	1.13	0.878742961
hTFIIS.h	AJ223473	0.98	1.14	0.856298131
HUNKI	Y12059	1.59	1.62	0.976707993
HUNKI	Y12059	1.33	1.50	0.884627755
HZF2	X78925	1.12	1.19	0.948487222
HZF2	X78925	1.08	1.19	0.908973223
HZF3	X78926	1.28	1.39	0.920945575
HZF3	X78926	1.10	1.31	0.838730175
HZF8	X78931	1.56	1.52	1.022134201
HZF8	X78931	1.40	1.56	0.896953681
HZF9	X78932	1.14	1.24	0.918602524
HZF9	X78932	1.11	1.30	0.857126824
Id1	NM_002165	1.24	1.23	1.00902126
Id1	NM_002165	1.13	1.41	0.80522294
Id3	A17548	1.38	1.31	1.055781754
Id3	X69111	1.27	1.28	0.990641606
Id4	Y07958	1.15	1.26	0.913664616
Id4	Y07958	1.09	1.32	0.830113526
INsAF	s73205	1.84	2.05	0.898920183
INsAF	s73205	1.85	2.13	0.871765981
intergenic	region U15407	2.30	2.60	0.88468389
HOXB7-HOXB6				
intergenic	region U15407	2.04	2.59	0.785453268
HOXB7-HOXB6				
IQGAP2	NM_006633	1.12	1.12	0.998484582
IQGAP2	NM_006633	0.94	1.12	0.840859025
IRF-1	X14454	2.41	2.57	0.938218115
IRF-1	X14454	2.39	2.58	0.925343204
IRF2	NM_002199	3.34	2.85	1.173965009
IRF2	NM_002199	2.94	2.56	1.14907375
IRF4	U52682	1.32	1.28	1.029933166
IRF4	U52682	1.37	1.43	0.959410817
IRF5	NM_002200	1.37	1.51	0.904052621
IRF5	NM_002200	1.36	1.59	0.858607001
IRF6	NM_006147	1.29	1.58	0.813425333
IRF6	NM_006147	1.18	1.50	0.789190299
IRF7	U53830	1.84	1.44	1.27973546
IRF7	NM_004029	1.32	1.21	1.084000454
Irx-4	NM_016358	1.19	1.15	1.029933166
Irx-4	NM_016358	1.17	1.22	0.956334448
IsGF-3gamma	M87503	1.42	1.55	0.915149715
IsGF-3gamma	M87503	1.39	1.56	0.887975373
Jun-D	X56681	2.38	2.25	1.056280294
Jun-D	X56681	2.04	2.18	0.933938896
JunB	X51345	1.02	1.14	0.892190868
JunB	X51345	0.98	1.14	0.855272625

K-ALPHA-1	NM_006082	0.83	0.96	0.86884485
K-ALPHA-1	NM_006082	0.83	0.97	0.859281424
KF1	NM_005667	0.93	1.05	0.890983333
KF1	NM_005667	0.91	1.06	0.864474263
KIAA0048	D28588	1.17	1.24	0.943988673
KIAA0048	D28588	1.19	1.30	0.918453567
KIAA0065	D31763	2.61	2.47	1.058679492
KIAA0065	D31763	2.53	2.52	1.005681703
KIAA0071	NM_015156	2.49	2.21	1.124047572
KIAA0071	NM_015156	2.30	2.27	1.015956269
KIAA0130	NM_014815	1.35	1.36	0.9886418
KIAA0130	NM_014815	1.17	1.34	0.869733568
KIAA0161	D79983	1.43	1.66	0.85937708
KIAA0161	D79983	1.42	1.69	0.837823111
KIAA0211	D86966	1.41	1.67	0.846204986
KIAA0211	D86966	1.37	1.73	0.79442123
KIAA0222	D86975	2.22	2.40	0.925360475
KIAA0222	D86975	2.02	2.43	0.82835128
KIAA0244	NM_015153	1.54	1.39	1.1095751
KIAA0244	NM_015153	1.45	1.36	1.067040755
KIAA0314	AB002312	2.38	2.57	0.927343337
KIAA0314	AB002312	2.33	2.65	0.876030662
KIAA0333	AB002331	1.05	1.22	0.861487483
KIAA0333	AB002331	1.07	1.25	0.854015656
KIAA0352	NM_014830	2.88	3.18	0.9057295
KIAA0352	NM_014830	2.37	2.80	0.8492877
KIAA0395	AB007855	1.56	1.77	0.879373168
KIAA0395	AB007855	1.42	1.77	0.801179995
KIAA0426	NM_014724	1.17	1.17	0.995781911
KIAA0426	NM_014724	1.06	1.23	0.866553199
KIAA0478	AB007947	2.27	2.38	0.954072874
KIAA0478	AB007947	2.25	2.62	0.857934327
KIAA0569	NM_014795	1.66	1.65	1.011174941
KIAA0569	NM_014795	1.39	1.76	0.78836631
KIAA0595	AB011167	1.90	1.85	1.026787356
KIAA0595	AB011167	1.71	2.19	0.782500333
KIAA0600	AB011172	1.90	1.34	1.413460448
KIAA0600	AB011172	2.18	1.58	1.381300612
KIAA0929	AB023146	1.54	1.62	0.949493335
KIAA0929	AB023146	1.55	1.63	0.949132006
KIAA1015	AB023232	2.58	2.62	0.982939758
KIAA1015	AB023232	2.17	2.68	0.811021231
KIAA1259	AB033085	0.85	1.04	0.817749165
KIAA1259	AB033085	0.91	1.18	0.771007682
KIAA1442	AB037863	2.14	2.34	0.914709549
KIAA1442	AB037863	2.15	2.39	0.898110711
KIAA1528	AB040961	6.42	6.40	1.003589265
KIAA1528	AB040961	6.67	6.98	0.955545485
KIAA1741	AW081989	1.58	1.79	0.882769399
KIAA1741	AW081989	1.68	1.99	0.846731464
KID	D38751	1.54	1.48	1.042612741
KID	D38751	1.45	1.52	0.95955196

KLF13	NM_015995	1.04	1.28	0.816419879
KLF13	NM_015995	0.91	1.14	0.796485569
KNSL4	AB017335	1.22	1.41	0.866676983
KNSL4	AB017335	1.19	1.45	0.818446687
Kox1	X52332	1.02	1.16	0.880125266
Kox1	X52332	0.98	1.24	0.789133958
Kox23	X52354	0.91	1.08	0.842330108
Kox23	X52354	0.90	1.08	0.832659332
Kox26	X52357	1.00	1.19	0.83622347
Kox26	X52357	0.99	1.26	0.785398373
Kox29	X52360	0.96	1.07	0.90087877
Kox29	X52360	0.98	1.09	0.897521031
Kox30	X52361	1.58	1.72	0.918425379
Kox30	X52361	1.38	1.53	0.902401118
KRAB	M67508	1.56	1.63	0.955633904
KRAB	M67508	1.47	1.60	0.922478172
Kruppel-type ZNF	AJ245587	2.04	2.40	0.851750841
Kruppel-type ZNF	AJ245587	1.79	2.14	0.836843037
KUP	X16576	0.96	1.15	0.839112982
KUP	X16576	0.92	1.13	0.816714046
L-Myc-1(long form)	X07262	1.05	1.20	0.876584744
L-Myc-1(long form)	X07262	1.01	1.23	0.826416004
LAF4	NM_002285	0.67	0.83	0.815483937
LAF4	NM_002285	0.65	0.84	0.784014372
LBR	NM_002296	1.25	1.30	0.966371608
LBR	NM_002296	1.23	1.38	0.891519857
LD5-1	U88080	1.15	1.38	0.82971758
LD5-1	U88080	1.11	1.41	0.790825308
LDOC1	NM_012317	1.33	1.41	0.946393907
LDOC1	NM_012317	1.28	1.42	0.897325005
LEF-1	AF203908	1.27	1.37	0.928294795
LEF-1	AF203908	1.16	1.44	0.810978586
lens epithelium-derived GF	AF063020	1.24	1.42	0.870186854
lens epithelium-derived GF	AF063020	1.13	1.39	0.81400662
leucine zipper	AF056184	2.24	2.72	0.824441293
leucine zipper	AF056184	2.47	3.10	0.796652442
leucine zipper kinase	AF251441	2.80	3.31	0.846204986
AZK	AF251441	2.71	3.35	0.808081689
LHX2	NM_004789	1.42	1.48	0.953866869
LHX2	NM_004789	1.33	1.52	0.87550605
LHX6	NM_014368	1.31	1.42	0.921284172
LHX6	NM_014368	1.28	1.42	0.905610681
LIM	AF061258	1.13	1.44	0.78655152
LIM	AF061258	1.09	1.41	0.773071315
LIM domain only 1 (rhombotin 1)	M26682	1.39	1.44	0.966564434
LIM domain only 1 (rhombotin 1)	M26682	1.32	1.49	0.883805842
LIM protein MLP	U49837	0.96	1.03	0.937706079
LIM protein MLP	U49837	0.95	1.14	0.82868354
LIM1	U14755	1.17	1.23	0.952256263
LIM1	U14755	1.01	1.27	0.798369687

LIMK	D26309	2.92	3.03	0.964180024
LIMK-2	D45906	1.60	1.66	0.965944664
LIMK-2	D45906	1.63	1.73	0.945399728
LMO4	U24576	0.85	0.84	1.007125772
LMO4	U24576	0.85	0.88	0.960803963
LOC51043	NM_015872	0.86	0.91	0.949928525
LOC51043	NM_015872	0.96	1.02	0.943797482
LOC51131	NM_016119	1.08	1.04	1.041898041
LOC51131	NM_016119	1.01	1.03	0.974830053
LOC51193	NM_016331	1.18	1.40	0.846337164
LOC51193	NM_016331	1.26	1.54	0.817829826
LOC51591	NM_015905	5.44	4.01	1.354983586
LOC51591	NM_015905	5.77	4.26	1.353823958
LOC51717	NM_016285	1.43	1.55	0.919576048
LOC51717	NM_016285	1.32	1.54	0.856030646
LOC55862	NM_018479	3.18	3.29	0.96566812
LOC55862	NM_018479	2.90	3.29	0.882904561
LOC56899	AF164792	1.35	1.46	0.923685155
LOC56899	AF164792	1.24	1.47	0.843151755
LyF-1	U40462	1.17	1.33	0.881798663
LyF-1	U40462	1.04	1.28	0.809448886
LZLP	NM_013344	1.63	1.78	0.914937922
LZLP	NM_013344	1.50	1.67	0.897409878
MADH4	NM_005359	1.48	1.23	1.202041185
MADH4	NM_005359	1.27	1.13	1.118588098
MADH5	NM_005903	1.19	1.37	0.867598314
MADH5	NM_005903	1.20	1.38	0.864484722
MAF	NM_005360	0.82	0.83	0.983383327
MAF	NM_005360	0.74	0.92	0.79706277
MAFG	NM_002359	1.33	1.60	0.833961234
MAFG	NM_002359	1.37	1.65	0.833526497
MAP4	NM_002375	3.80	4.62	0.824293011
MAP4	NM_002375	3.69	4.71	0.78417244
MAPK8	NM_002750	0.88	1.00	0.88152506
MAPK8	NM_002750	0.88	1.02	0.860049569
MAZ	M94046	1.21	1.47	0.819442731
MAZ	M94046	1.19	1.48	0.804052549
MB67	Z30425	1.08	1.02	1.060408157
MB67	Z30425	0.99	1.08	0.915952791
MCG4	NM_006782	1.15	1.31	0.87362439
MCG4	NM_006782	1.15	1.34	0.857557298
MEF2A	U49020	1.18	1.29	0.917750293
MEF2A	U49020	1.08	1.27	0.851735738
MEF2B	NM_005919	1.02	1.07	0.950137026
MEF2B	NM_005919	0.95	1.04	0.910069513
MEF2D	NM_005920	1.39	1.33	1.043108425
MEF2D	NM_005920	1.20	1.44	0.837034123
metallopanstimulin	U85979	1.98	2.01	0.985678226
metallopanstimulin	U85979	1.94	2.20	0.882570172
MHox (K-2)	M95929	1.07	1.17	0.914292266
MHox (K-2)	M95929	0.95	1.17	0.810474232
Mi	Z29678	1.71	1.66	1.030471845

Mi	Z29678	1.76	1.79	0.986205176
MITF	AF034755	1.23	1.24	0.99130983
MITF	AF034755	1.32	1.50	0.883057308
Miz-1	Y09723	1.01	1.15	0.876000186
Miz-1	Y09723	0.93	1.14	0.814161592
MLH3	NM_005784	0.54	0.63	0.855999025
MLH3	NM_005784	0.62	0.78	0.801334083
MLX	AF203978	1.41	1.49	0.949042398
MLX	AF203978	1.36	1.48	0.923306997
Mog	U64564	1.32	1.37	0.960484338
Mog	U64564	1.27	1.39	0.915925265
MRG1	AF109161	3.76	4.37	0.860312626
MRG1	AF109161	3.73	4.50	0.827783082
MTERF	NM_006980	1.51	1.80	0.838119573
MTERF	NM_006980	1.35	1.70	0.789625228
MTF-1	AJ251881	2.11	2.39	0.881959401
MTF-1	AJ251881	1.95	2.39	0.815353763
mtTF1	X64269	1.47	1.59	0.925536704
mtTF1	X64269	1.44	1.57	0.914838473
MXI1	NM_005962	1.16	1.29	0.898657286
MXI1	NM_005962	1.16	1.36	0.857867078
MYBBP1A	AF147709	2.29	1.77	1.292847997
MYBBP1A	AF147709	1.85	1.75	1.054649057
MYCBP	NM_012333	3.73	3.58	1.040887845
MYCBP	NM_012333	3.47	3.48	0.997884909
MYCL2	NM_005377	2.12	2.03	1.044677307
MYCL2	NM_005377	2.04	2.00	1.018897998
MYCLK1	M64786	1.43	1.73	0.828883125
MYCLK1	M64786	1.49	1.80	0.826354974
MYT2	NM_003871	4.01	4.17	0.962205771
MYT2	NM_003871	4.04	4.42	0.915182881
N-CoR	AF044209	1.33	1.29	1.027153581
N-CoR	AF044209	1.25	1.29	0.969389141
N-Oct-3	Z11933	3.50	3.17	1.103689021
N-Oct-3	Z11933	2.91	3.05	0.955346496
N143	AJ002572	3.89	3.16	1.232431216
N143	AJ002572	2.82	3.41	0.828068155
NACA	NM_005594	1.34	1.26	1.061449635
NACA	NM_005594	1.22	1.36	0.899257451
NAGA	NM_000262	2.23	2.55	0.873072079
NAGA	NM_000262	2.02	2.54	0.795967326
NCOA1	NM_003743	1.34	1.43	0.939022342
NCOA1	NM_003743	1.36	1.45	0.932646647
NCOA3	NM_006534	2.14	2.15	0.995002762
NCOA3	NM_006534	1.97	2.05	0.959254041
NCYM	NM_006316	1.18	1.11	1.067219564
NCYM	NM_006316	1.07	1.16	0.917384574
NDUFA6	NM_002490	0.80	0.82	0.969899497
NDUFA6	NM_002490	0.71	0.92	0.772339515
Negative control	Negative control	1.29	1.11	1.161392449
Negative control	Negative control	5.43	5.29	1.027043989
NEUROD2	U58681	1.14	1.28	0.889551897

NEUROD2	U58681	1.02	1.28	0.795592113
NEUROG1	U63842	1.39	1.71	0.812574039
NEUROG1	U63842	1.29	1.63	0.795487149
NF-1X	U07811	0.99	0.82	1.215806558
NF-1X	U07811	0.64	0.82	0.782275487
NFAT1	U43341	2.28	2.65	0.861852199
NFAT1	U43341	2.30	2.80	0.819721245
NFATC1	NM_006162	1.21	1.27	0.956885723
NFATC1	NM_006162	1.20	1.33	0.906442678
NFATX	U14510	1.09	1.35	0.8066644
NFATX	U14510	0.99	1.24	0.798995238
NFIL3	NM_005384	3.33	3.43	0.969982487
NFIL3	NM_005384	3.22	3.36	0.957589194
NFKB1	M58603	2.44	2.68	0.910234175
NFKB1	M55643	1.23	1.37	0.894069494
NFKB2	U09609	1.09	1.21	0.899003953
NFKB2	U09609	1.04	1.28	0.815426014
NFKBIB	NM_002503	0.66	0.74	0.891632657
NFKBIB	NM_002503	0.61	0.73	0.835589511
NFKBIE	NM_004556	1.34	1.34	0.995844935
NFKBIE	NM_004556	1.30	1.37	0.951450999
NFkBp105	M55643	0.82	0.80	1.028950235
NFkBp105	M55643	0.78	0.88	0.882630106
ngn3	AJ133776	1.04	1.14	0.911675496
ngn3	AJ133776	1.01	1.15	0.877939013
NME2	NM_002512	1.08	1.28	0.849242645
NME2	NM_002512	0.99	1.21	0.819907039
Nmi	U32849	1.50	1.65	0.908824603
Nmi	U32849	1.46	1.67	0.874987469
NOD1	AF149774	0.88	0.97	0.913819737
NOD1	AF149774	0.80	0.97	0.829329103
NOT3	NM_014516	1.10	1.05	1.043010425
NOT3	NM_014516	1.01	1.32	0.770565768
NP220	D83032	1.59	1.80	0.886060234
NP220	D83032	1.50	1.74	0.861556367
NPAS1	NM_002517	2.55	3.06	0.832115639
NPAS1	NM_002517	2.53	3.18	0.795687598
NR0B1	NM_000475	0.89	0.90	0.985394227
NR0B1	NM_000475	0.84	1.02	0.822398708
NR2F6	NM_005234	1.11	1.26	0.878406569
NR2F6	NM_005234	1.03	1.23	0.843593242
NR3C1	NM_000176	1.45	1.63	0.884627755
NR3C1	NM_000176	1.37	1.57	0.872392013
NR4A2	NM_006186	5.15	5.52	0.933470433
NR4A2	NM_006186	4.88	5.68	0.859257687
NR5A1	NM_004959	1.55	1.88	0.827098402
NR5A1	NM_004959	1.56	1.94	0.806250074
NRL	M81840	1.12	1.36	0.824507422
NRL	M81840	1.08	1.34	0.802675923
NRsF form 2	U13879	1.51	1.60	0.940689035
NRsF form 2	U13879	1.29	1.56	0.826356612
NSEP1	NM_004559	4.08	4.54	0.898882683

NSEP1	NM_004559	4.09	4.85	0.843016697
nuclear factor 1 B-type	U07810	1.67	1.72	0.970651102
nuclear factor 1 B-type	U07810	1.50	1.56	0.957498055
nuclear factor I-B2	U85193	5.86	6.80	0.86191166
nuclear factor I-B2	U85193	6.04	7.09	0.85215085
nuclear factor IV	X57500	1.44	1.28	1.118113871
nuclear factor IV	X57500	1.35	1.53	0.882482444
OAZ	AF221712	0.95	0.98	0.974744849
OAZ	AF221712	0.82	0.94	0.867200363
Oct-1B=POU homeodomain	S66902	1.11	1.23	0.901795827
Oct-1B=POU homeodomain	S66902	1.07	1.22	0.879755632
Oct-4A	Z11900	1.44	1.75	0.825538314
Oct-4A	Z11900	1.43	1.85	0.773109431
OGL12	AF023203	2.31	2.68	0.864811744
OGL12	AF023203	2.45	2.92	0.837215348
OSMRB	U60805	0.90	1.09	0.825581
OSMRB	U60805	0.78	0.97	0.805457565
OTF3C	Z11901	6.19	4.42	1.40097838
OTF3C	Z11901	5.89	4.42	1.332945525
OTX1	AB037501	1.87	1.84	1.018457389
OTX1	AB037501	1.75	1.86	0.938379
OVOL1	NM_004561	1.34	1.20	1.12018921
OVOL1	NM_004561	1.05	1.21	0.869990813
p130	s67171	2.11	1.71	1.231079229
p130	s67171	1.56	1.60	0.975948711
p243	AJ242977	1.27	1.43	0.884227005
p243	AJ242977	1.37	1.69	0.812266478
P38IP	NM_017569	0.99	1.09	0.90720651
P38IP	NM_017569	0.94	1.10	0.856087428
p53	K03199	1.39	1.68	0.831158406
p53	K03199	1.38	1.67	0.828166161
p621	AJ242978	1.19	1.20	0.990516633
p621	AJ242978	1.06	1.18	0.898178687
PACE4	NM_002570	1.29	1.48	0.867988727
PACE4	NM_002570	1.26	1.48	0.852795758
PAX1	NM_006192	1.20	1.32	0.903818349
PAX1	NM_006192	1.06	1.32	0.807894213
PAX2	U45255	1.46	1.62	0.901165711
PAX2	U45255	1.40	1.63	0.858751434
PAX3	NM_000438	1.21	1.36	0.886358667
PAX3	NM_000438	1.16	1.37	0.850478749
PAX5	U56835	0.93	1.06	0.871188843
PAX5	NM_016734	1.71	2.02	0.846266
PAX6	U63833	1.33	1.53	0.865756264
PAX6	U63833	1.27	1.52	0.833925287
PAX8	S55490	1.94	2.07	0.934433059
PAX8	S55490	1.82	2.05	0.885815601
PAX9	NM_006194	0.78	0.95	0.817959194
PAX9	X92850	1.17	1.49	0.784900153
PBX1	NM_002585	1.46	1.26	1.160624187
PBX1	NM_002585	1.56	1.36	1.14486291
PBX2	NM_002586	1.14	1.28	0.885932245

PBX2	NM_002586	1.14	1.34	0.848773413
PC4	NM_006713	0.70	0.80	0.879994421
PC4	NM_006713	0.70	0.81	0.86301099
PCAF	NM_003884	1.07	1.34	0.798415415
PCAF	NM_003884	1.03	1.29	0.796481403
PDEF	NM_012391	1.17	1.33	0.876561702
PDEF	NM_012391	1.38	1.70	0.811137032
PEA3	D12765	1.21	1.54	0.784602478
PEA3	D12765	1.21	1.56	0.775569587
PEPD	J04605	0.66	0.78	0.8570049
PEPD	J04605	0.71	0.86	0.827558815
PGF	NM_002632	1.08	1.18	0.917061973
PGF	NM_002632	1.01	1.20	0.83593402
pGLI3HH	M20674	1.21	1.31	0.921799298
pGLI3HH	M20674	1.13	1.32	0.856812924
PIAS3	NM_006099	3.47	4.09	0.849170482
PIAS3	NM_006099	3.59	4.26	0.842175439
PINCH	U09284	1.62	1.49	1.088214315
PINCH	U09284	1.44	1.44	1.001116263
Pit-1	D10216	2.56	2.74	0.934572932
Pit-1	D10216	2.26	2.72	0.831778211
PITX1	NM_002653	1.04	1.23	0.841903944
PITX1	NM_002653	0.95	1.22	0.780247958
PITX2	U69961	2.17	1.90	1.142972468
PITX2	U69961	1.90	1.73	1.099447256
PITX3	NM_005029	1.08	1.19	0.908904038
PITX3	NM_005029	1.05	1.16	0.900257494
PKNOX1	NM_004571	2.66	2.91	0.915727678
PKNOX1	NM_004571	2.43	2.80	0.867326045
PLCG1	NM_002660	0.88	1.09	0.801330479
PLCG1	NM_002660	0.87	1.12	0.777390419
PML	M79462	2.93	3.22	0.90918611
PML	M79462	2.83	3.18	0.891376967
POU6F1	NM_002702	1.18	1.38	0.853348217
POU6F1	NM_002702	1.11	1.39	0.803725887
PPAR delta	AF187850	1.68	2.05	0.817974153
PPAR delta	AF187850	1.63	2.05	0.797208472
PPARbeta	L07592	1.12	1.30	0.860735779
PPARbeta	L07592	1.09	1.30	0.841097109
PPARBP	NM_004774	2.49	2.42	1.028888699
PPARBP	NM_004774	2.59	2.62	0.989587372
PPARG	NM_005037	1.76	1.92	0.919451555
PPARG	NM_005037	1.54	1.87	0.82431469
PPARGC1	NM_013261	4.02	4.08	0.985608174
PPARGC1	NM_013261	3.61	3.97	0.910308604
PPIH	NM_006347	1.11	1.36	0.810066673
PPIH	NM_006347	1.10	1.37	0.797849135
pRb	X16439	1.29	1.40	0.923240454
pRb	X16439	1.21	1.44	0.839720657
PRDM4	NM_012406	1.09	1.14	0.952491603
PRDM4	NM_012406	1.02	1.09	0.935085892
protein Id4	U28368	1.18	1.11	1.06450375

protein Id4	U28368	1.29	1.23	1.054319434
protein p38	AJ242975	1.63	1.95	0.834950074
protein p38	AJ242975	1.44	1.85	0.781801717
PRX2	NM_016307	4.82	3.63	1.326456916
PRX2	NM_016307	4.32	4.25	1.017671923
PSCDBP	NM_004288	0.74	0.85	0.86992699
PSCDBP	NM_004288	0.69	0.84	0.81936229
PSMC1	NM_002802	1.36	1.52	0.894759062
PSMC1	NM_002802	1.18	1.33	0.891456342
PTHR1	NM_000316	1.31	1.42	0.924499528
PTHR1	NM_000316	1.20	1.40	0.8556512
PXMP3	NM_000318	1.62	2.02	0.804068402
PXMP3	NM_000318	1.40	1.80	0.78066981
PXN	NM_002859	2.72	2.90	0.93925013
PXN	NM_002859	2.51	2.90	0.863012935
rab 13	X75593	1.25	1.23	1.008775651
rab 13	X75593	1.12	1.33	0.83974611
RAR-alpha1	X06614	1.43	1.62	0.88166305
RAR-alpha1	X06614	1.30	1.60	0.814209521
RAR-b	M96016	1.57	1.95	0.801619789
RAR-b	M96016	1.57	2.00	0.782949951
RARA	NM_000964	1.42	1.65	0.862685131
RARA	NM_000964	1.40	1.65	0.847918555
RARG	NM_000966	1.42	1.61	0.882296859
RARG	NM_000966	1.41	1.60	0.882261145
RB1	NM_000321	0.97	1.19	0.812728745
RB1	NM_000321	0.96	1.20	0.800478062
RBL1	NM_002895	2.04	2.42	0.841470759
RBL1	NM_002895	1.92	2.35	0.817621225
RBP-L	AB026048	0.94	0.70	1.339824561
RBP-L	AB026048	0.80	0.71	1.133824475
RCL	NM_006443	1.26	1.39	0.906711617
RCL	NM_006443	1.24	1.39	0.891529469
RELA	Z22951	0.86	0.89	0.96567493
RELA	Z22951	0.80	0.85	0.94852389
repressor protein	D30612	1.37	1.53	0.890929984
repressor protein	D30612	1.32	1.52	0.87316143
REQ	NM_006268	1.43	1.67	0.860238236
REQ	NM_006268	1.46	1.72	0.847602428
retinoid X receptor	U66306	2.17	2.44	0.889382828
retinoid X receptor	U38480	2.07	2.33	0.889199662
gamma RFP	NM_006510	3.73	3.97	0.940382915
RFP	NM_006510	3.81	4.43	0.858648769
RFX3	X76092	1.62	1.47	1.1024024
RFX3	X76092	1.43	1.62	0.8816817
rhoHP1	D85815	1.42	1.33	1.069393174
rhoHP1	D85815	1.46	1.53	0.955107329
RING1	NM_002931	1.42	1.59	0.896164283
RING1	NM_002931	1.41	1.60	0.880810591
RLF	NM_012421	3.38	3.75	0.901013305
RLF	NM_012421	3.65	4.07	0.898102049
RNF NY-REN-43	AF155109	1.18	1.29	0.913146151

RNF NY-REN-43	AF155109	1.16	1.43	0.811644103
RNF13	NM_007282	1.22	1.33	0.916119358
RNF13	NM_007282	1.20	1.31	0.912867135
RNF15	NM_006355	1.29	1.45	0.893216374
RNF15	NM_006355	1.17	1.44	0.811128245
RNF4	NM_002938	1.35	1.44	0.936370857
RNF4	NM_002938	1.32	1.45	0.907740218
RNF9	NM_006778	1.25	1.36	0.918123369
RNF9	NM_006778	1.18	1.36	0.863185084
RNP-specific A	X06347	1.31	1.39	0.94551044
RNP-specific A	X06347	1.16	1.47	0.788038353
RORalpha2	U04898	4.15	4.42	0.938764319
RORalpha2	U04898	4.03	4.29	0.938708029
RORbeta	Y08639	1.29	1.50	0.858111801
RORbeta	Y08639	1.27	1.50	0.842642276
RORC	NM_005060	1.39	1.61	0.861315789
RORC	NM_005060	1.43	1.77	0.807520338
RP58	AJ223321	1.34	1.40	0.953320654
RP58	AJ223321	1.19	1.38	0.866072097
RPF-1	U91934	1.26	1.51	0.833565324
RPF-1	U91934	1.23	1.50	0.822227125
RPL13A	X56932	0.87	0.88	0.991870123
RPL13A	X56932	0.77	0.87	0.883814097
RPL15	NM_002948	1.01	1.07	0.944600915
RPL15	NM_002948	0.98	1.14	0.859452181
RPL21	NM_000982	1.60	1.53	1.04809166
RPL21	NM_000982	1.57	1.58	0.995425213
RPL23A	NM_000984	1.59	1.42	1.117137899
RPL23A	NM_000984	1.46	1.38	1.059317332
RPL37	NM_000997	1.09	1.23	0.883744302
RPL37	NM_000997	1.09	1.30	0.842639916
RPS11	NM_001015	1.55	1.32	1.171184602
RPS11	NM_001015	1.30	1.22	1.068789518
RPS19	NM_001022	0.84	1.00	0.841774594
RPS19	NM_001022	0.86	1.05	0.819892642
RRN3	NM_018427	1.64	1.61	1.015843155
RRN3	NM_018427	1.10	1.40	0.78552954
RUVBL1	NM_003707	1.21	1.41	0.864080705
RUVBL1	NM_003707	1.15	1.43	0.805614035
Rx	AF001911	1.40	1.19	1.169408279
Rx	AF001911	1.21	1.29	0.940515001
RXR-alpha	X52773	1.14	1.20	0.95178794
RXR-alpha	X52773	1.02	1.17	0.875212013
RXRB	U00961	1.41	1.76	0.802764083
RXRB	U00961	1.32	1.64	0.802187954
SAFB	NM_002967	2.08	1.85	1.122521568
SAFB	NM_002967	1.98	1.85	1.072239203
SALL1	NM_002968	1.06	1.32	0.799379966
SALL1	NM_002968	1.09	1.37	0.794919835
sAP-1a	M85165	1.02	1.15	0.893216374
sAP-1a	M85165	0.99	1.14	0.868660598
SEP3B	AF285109	1.36	1.52	0.895524427

SEP3B	AF285109	1.34	1.51	0.891662954
sF1	D88155	1.24	1.23	1.006655807
sF1	D88155	0.89	1.10	0.815406356
SF3A1	NM_005877	0.94	1.18	0.796947498
SF3A1	NM_005877	0.98	1.24	0.789354005
SIX1	X91868	1.28	1.26	1.011940877
SIX1	X91868	1.15	1.27	0.899649002
SIX6	AF141651	1.31	1.51	0.866808238
SIX6	AF141651	1.28	1.61	0.795100662
SKI	NM_003036	1.27	1.34	0.951830965
SKI	NM_003036	1.23	1.34	0.916814067
SKIL	NM_005414	1.22	1.23	0.996060051
SKIL	NM_005414	1.18	1.23	0.965665088
Smad2	U78726	1.54	1.70	0.902843033
Smad2	U78726	1.52	1.74	0.876401307
SMARCA3	NM_003071	2.60	2.63	0.988267744
SMARCA3	NM_003071	2.52	2.58	0.977550509
SMARCA4	NM_003072	1.20	1.41	0.850958457
SMARCA4	NM_003072	1.14	1.43	0.798506046
SMARCC1	NM_003074	1.37	1.53	0.897049921
SMARCC1	NM_003074	1.31	1.51	0.867695906
SMARCC2	NM_003075	1.11	1.36	0.816630385
SMARCC2	NM_003075	1.10	1.37	0.803295263
SMN1	U18423	2.14	2.06	1.0410434
SMN1	U18423	1.93	2.06	0.938075938
SNAP190	AF032387	1.08	1.15	0.940066948
SNAP190	AF032387	1.19	1.34	0.88972042
SNAPC3	NM_003084	0.64	0.65	0.973605848
SNAPC3	NM_003084	0.58	0.67	0.873988625
snRNP B	X17567	0.99	0.98	1.010085806
snRNP B	X17567	0.82	0.97	0.840177885
SOX10	AJ001183	1.71	1.82	0.937524016
SOX10	AJ001183	1.59	1.78	0.894532832
SOX13	NM_005686	1.71	1.92	0.891384762
SOX13	NM_005686	1.66	1.93	0.860515571
SOX4	X70683	0.90	0.93	0.960960313
SOX4	X70683	0.82	0.92	0.894488762
SOX6	X65663	0.69	0.79	0.882795517
SOX6	X65663	0.65	0.75	0.87280897
SOX8	AF164104	1.79	2.09	0.857375717
SOX8	AF164104	1.65	2.13	0.774778844
SOX9	Z46629	1.69	1.88	0.898185589
SOX9	Z46629	1.55	1.92	0.807916038
SP1	J03133	1.18	1.30	0.909375062
SP1	J03133	1.16	1.30	0.887824726
SP3	X68560	1.66	1.66	1.001079125
SP3	X68560	1.45	1.77	0.818395433
sRF	J03161	1.45	1.67	0.867315899
sRF	J03161	1.43	1.69	0.84824421
sRY	L10101	1.18	1.21	0.982441028
sRY	L10101	1.13	1.21	0.934783803
STAT2	M97934	1.41	1.52	0.926980567

STAT2	M97934	1.41	1.62	0.868724958
STAT5B	NM_012448	1.56	1.40	1.114847778
STAT5B	NM_012448	1.40	1.71	0.821978259
STAT6	NM_003153	1.27	1.36	0.938049629
STAT6	NM_003153	1.21	1.37	0.879605458
SZF1	NM_016089	1.21	1.50	0.802961061
SZF1	NM_016089	1.16	1.49	0.774800507
T-STAR	NM_006558	0.67	0.78	0.861849244
T-STAR	NM_006558	0.68	0.82	0.831664605
T3R	Y00479	1.71	1.62	1.053389262
T3R	Y00479	1.60	1.53	1.041322337
T3R	X55066	1.56	1.89	0.824459774
TAF(I)63	L39061	1.00	1.12	0.896350467
TAF(I)63	L39061	1.02	1.30	0.783696377
TAF(II)30	U25816	1.51	1.40	1.074026032
TAF(II)30	U25816	1.40	1.38	1.015134059
TAF(II)32	U21858	0.98	1.22	0.802461769
TAF(II)32	U21858	0.98	1.23	0.792812413
TAF(II)70-alpha	L25444	0.96	1.02	0.941312641
TAF(II)70-alpha	L25444	0.90	1.03	0.872787029
TAF2A	NM_004606	1.13	1.14	0.999330892
TAF2A	NM_004606	0.93	1.11	0.838249213
TAF2F	NM_005642	1.03	1.26	0.81530342
TAF2F	NM_005642	1.02	1.30	0.784947723
TAF2I	NM_005643	1.50	1.43	1.045218429
TAF2I	NM_005643	1.39	1.41	0.991353741
TAF2I	AF118094	1.11	1.26	0.881120736
TAF2J	NM_005644	1.28	1.40	0.913232427
TAF2J	NM_005644	1.23	1.45	0.849684168
TAF2K	NM_005645	2.39	2.40	0.997067405
TAF2K	NM_005645	2.40	2.47	0.972697492
TAFII105	Y09321	1.26	1.45	0.867396208
TAFII105	Y09321	1.19	1.45	0.818176516
Tal-1	NM_003189	1.37	1.51	0.902488517
Tal-1	NM_003189	1.27	1.53	0.828923165
TARBP2	NM_004178	1.09	1.24	0.873471591
TARBP2	NM_004178	1.08	1.37	0.786576406
TBP	NM_003194	2.77	2.67	1.040623399
TBP	NM_003194	2.51	2.58	0.973841382
TBPL1	NM_004865	1.28	1.30	0.987855849
TBPL1	NM_004865	1.11	1.40	0.794804113
TBR1	NM_006593	1.19	1.20	0.996692807
TBR1	NM_006593	1.05	1.26	0.836636729
TBX19	NM_005149	1.36	1.48	0.923280344
TBX19	NM_005149	1.44	1.59	0.909429873
TBX2	NM_005994	0.85	1.07	0.798962867
TBX2	NM_005994	0.83	1.06	0.785853316
TBX20	AJ237589	1.34	1.21	1.102533818
TBX20	AJ237589	1.36	1.64	0.831910222
TBX6	NM_004608	4.15	4.62	0.899551242
TBX6	NM_004608	3.96	4.53	0.875140298
TCEA1	NM_006756	1.27	1.14	1.110125734

TCEA1	NM_006756	1.16	1.11	1.05196336
TCEB2	NM_007108	0.58	0.43	1.337134151
TCEB2	NM_007108	0.51	0.40	1.277713489
TCF-1	Z47365	1.00	1.16	0.86404602
TCF-1	Z47365	0.92	1.18	0.774647829
TCF-4	Y11306	1.36	1.46	0.928464375
TCF-4	Y11306	1.32	1.56	0.849608167
TCF21	NM_003206	1.75	2.02	0.863693344
TCF21	NM_003206	1.69	2.12	0.798699048
TCF4	NM_003199	0.93	0.92	1.021001263
TCF4	NM_003199	0.84	0.92	0.912194059
TCF6L1	NM_003201	1.67	1.95	0.857799865
TCF6L1	NM_003201	1.82	2.34	0.776883554
TCFL1	NM_005997	1.45	1.63	0.891060583
TCFL1	NM_005997	1.45	1.68	0.863907712
TCFL5	NM_006602	1.87	2.31	0.809256058
TCFL5	NM_006602	1.79	2.27	0.788010751
TEAD1	M63896	1.97	2.40	0.821174945
TEAD1	M63896	1.84	2.35	0.783305005
TEF-4	X94440	1.14	1.29	0.883210896
TEF-4	X94440	1.13	1.33	0.854457478
TF	U79243	1.42	1.54	0.919800195
TF	U79243	1.29	1.63	0.789904601
TFCP2	NM_005653	0.99	1.11	0.887949768
TFCP2	NM_005653	0.94	1.13	0.832221275
TFE3	AL161985	1.20	1.25	0.952888449
TFE3	AL161985	1.16	1.30	0.896818053
TFIIA	NM_015859	0.84	0.82	1.018380452
TFIIA	NM_015859	0.80	0.82	0.977671128
TFIID	Z22828	2.50	2.34	1.068758898
TFIID	Z22828	2.55	2.73	0.936976254
TFIIH-cyclin H	U11791	1.28	0.95	1.34843684
TFIIH-cyclin H	U11791	1.29	0.98	1.318842969
TFIIH-MO15	X77743	2.43	2.39	1.014330459
TFIIH-MO15	X77743	2.43	2.39	1.012865369
TFIIH-p34	Z30093	2.74	2.96	0.92722107
TFIIH-p34	Z30093	2.34	2.95	0.792209645
TFRC	NM_003234	1.33	1.49	0.895571075
TFRC	NM_003234	1.28	1.59	0.809200973
TGIF	NM_003244	1.75	1.38	1.274809288
TGIF	NM_003244	1.52	1.51	1.004868421
TIEG2	NM_003597	1.39	1.48	0.938352308
TIEG2	NM_003597	1.35	1.51	0.889349637
TIF1GAMMA	NM_015906	1.01	1.27	0.800846532
TIF1GAMMA	NM_015906	1.01	1.30	0.777179202
TIF2	X97674	1.33	1.45	0.922316636
TIF2	X97674	1.31	1.45	0.902108121
TIM44	NM_006351	1.33	1.57	0.847735407
TIM44	NM_006351	1.36	1.61	0.84323103
Timeless	AF098162	2.00	2.39	0.833636058
Timeless	AF098162	1.93	2.42	0.796169622
TIMM8b	AF152350	1.40	1.42	0.984849477

TIMM8b	AF152350	1.42	1.51	0.941288783
TIMM9	NM_012460	0.71	0.77	0.920513309
TIMM9	NM_012460	0.68	0.84	0.805700618
Tis11d	U07802	1.12	1.35	0.827564107
Tis11d	U07802	1.04	1.27	0.821167897
TNRC11	NM_005120	0.88	1.05	0.836486567
TNRC11	NM_005120	0.85	1.05	0.802230359
TOB1	NM_005749	1.02	1.29	0.790953507
TOB1	NM_005749	1.02	1.32	0.77856472
TOP1	U07806	3.49	3.40	1.026563028
TOP1	U07806	3.18	3.15	1.008577791
TP53BP1	NM_005657	0.83	0.86	0.969466553
TP53BP1	NM_005657	0.79	0.86	0.910836728
TP73	NM_005427	4.36	4.73	0.923146915
TP73	NM_005427	3.95	4.55	0.867754692
TR2	AF171055	1.60	1.59	1.007806633
TR2	AF171055	1.53	1.69	0.90288023
TRAF6	NM_004620	1.25	1.38	0.902520712
TRAF6	NM_004620	1.33	1.62	0.825230675
TTF-1	U43203	1.57	1.92	0.818730798
TTF-1	U43203	1.59	1.97	0.8042506
TTF-I	interacting AF000560	0.93	1.01	0.912809508
peptide				
TTF-I	interacting AF000560	0.92	1.02	0.908180051
peptide				
TTF1	NM_007344	1.36	1.32	1.03206288
TTF1	NM_007344	1.21	1.33	0.908191402
TTP	M63625	1.47	1.69	0.871059069
TTP	M63625	1.45	1.78	0.812551459
tumor suppressor	AJ224819	0.97	0.96	1.010680445
tumor suppressor	AJ224819	0.94	0.94	1.002997158
twist	X91662	1.15	1.30	0.889607419
twist	X91662	1.14	1.32	0.862171118
TZFP	NM_014383	1.61	1.57	1.026900096
TZFP	NM_014383	1.31	1.62	0.806125978
ubiquitin	M26880	1.25	1.29	0.968558812
ubiquitin	M26880	1.20	1.38	0.866123555
UBP1	NM_014517	1.16	1.39	0.837548498
UBP1	NM_014517	1.05	1.31	0.801834157
UKLF	AB015132	0.94	1.15	0.812199016
UKLF	AB015132	0.91	1.13	0.807110731
UsF1	X55666	0.92	0.77	1.202641159
UsF1	X55666	0.90	1.16	0.779296001
UsF2	X90824	1.70	1.51	1.12444728
UsF2	X90824	1.49	1.47	1.010033818
UTF1	NM_003577	0.78	0.92	0.852557876
UTF1	NM_003577	0.75	0.88	0.846901451
Vax-2	Y17791	1.95	1.73	1.125466134
Vax-2	Y17791	1.50	1.58	0.944890049
VDR	NM_000376	2.14	1.94	1.102535767
VDR	NM_000376	2.17	1.98	1.096166462
Vimentin	X56134	0.85	0.82	1.03779146
Vimentin	X56134	0.77	0.78	0.995470101
VSX1	NM_014588	1.19	1.38	0.862794625

VSX1	NM_014588	1.14	1.36	0.838036984
WAVE2	AB026542	1.37	1.57	0.873152446
WAVE2	AB026542	1.34	1.56	0.8602453
Whn	Y11746	0.95	1.05	0.89877812
Whn	Y11739	0.98	1.10	0.8889781
winged-helix	AF055080	1.80	1.62	1.112375194
TFforkhead 5	AF055080	1.64	1.65	0.995925632
winged-helix	AF055080	1.64	1.65	0.995925632
TFforkhead 5	AF055080	1.64	1.65	0.995925632
XB	U52701	0.93	1.00	0.931696975
XB	U52701	0.86	0.99	0.874287286
XBP1	NM_005080	1.32	1.48	0.894006132
XBP1	NM_005080	1.29	1.50	0.861985033
XG	Z48514	0.94	1.07	0.879021004
XG	Z48514	0.94	1.11	0.844934408
XPE-BF	U32986	1.23	1.06	1.157546744
XPE-BF	U32986	1.11	1.18	0.939655721
XPOT	NM_007235	0.93	1.04	0.888739099
XPOT	NM_007235	0.91	1.09	0.833312043
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YAF2	U72209	1.29	1.55	0.831250433
YPT3	X79780	1.04	1.04	0.999551657
YPT3	X79780	0.91	1.09	0.841560488
YWHAZ	NM_003406	1.42	1.48	0.955552146
YWHAZ	NM_003406	1.34	1.45	0.924713154
ZFD25	AB027251	1.38	1.47	0.935259419
ZFD25	AB027251	1.38	1.59	0.867549237
ZFM1	D26120	1.38	1.52	0.904756638
ZFM1	D26120	1.32	1.51	0.870056053
ZFN3	X60153	1.11	1.27	0.873020321
ZFN3	X60153	1.08	1.27	0.84639825
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ZFP37	NM_003408	2.85	3.53	0.806477053
ZFP37	NM_003408	3.00	3.78	0.795656333
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ZFS-2	D70832	1.19	1.34	0.887098454
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zinc finger factor GKLF	AF105036	2.21	2.60	0.850960542
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ZK1	NM_005815	1.13	1.34	0.848599298
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ZNF	AF024708	0.85	1.06	0.803894737
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ZNF	AF244088	0.96	1.18	0.80824225

ZNF	AL359576	1.96	2.40	0.816139265
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ZNF	L14787	0.77	0.99	0.776247563
ZNF	L14843	1.04	1.21	0.859382421
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ZNK75a	X91826	0.98	1.24	0.792918773
ZRP-1	AF000974	4.16	4.74	0.87742034
ZRP-1	AF000974	4.06	4.99	0.813590373
ZYX	NM_003461	1.40	1.36	1.031031823
ZYX	NM_003461	1.31	1.36	0.963983275

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